

Scheme 1

Solid Phase Protein Synthesis

Native Chemical Ligations in an N- to C-Terminal Direction

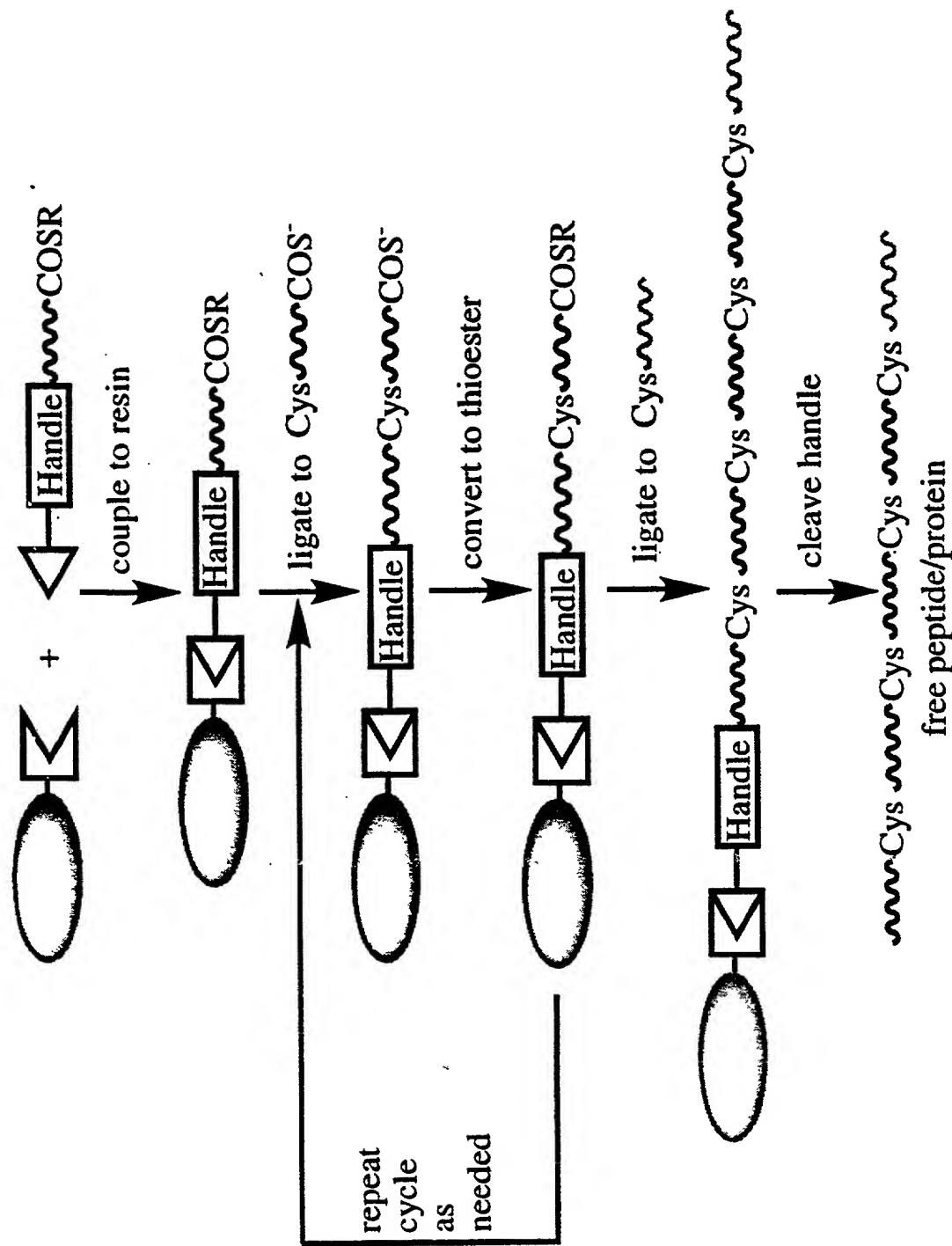


FIG. 1

Cys + COSY Stability Under Ligation Conditions

1. In the absence of a thioester peptide

H-CGFRVREFGDNTA-COSH MW=1487.6

6M GU·HCL, 0.1M NaPi, 0.5% thiophenol, room temperature, overnight

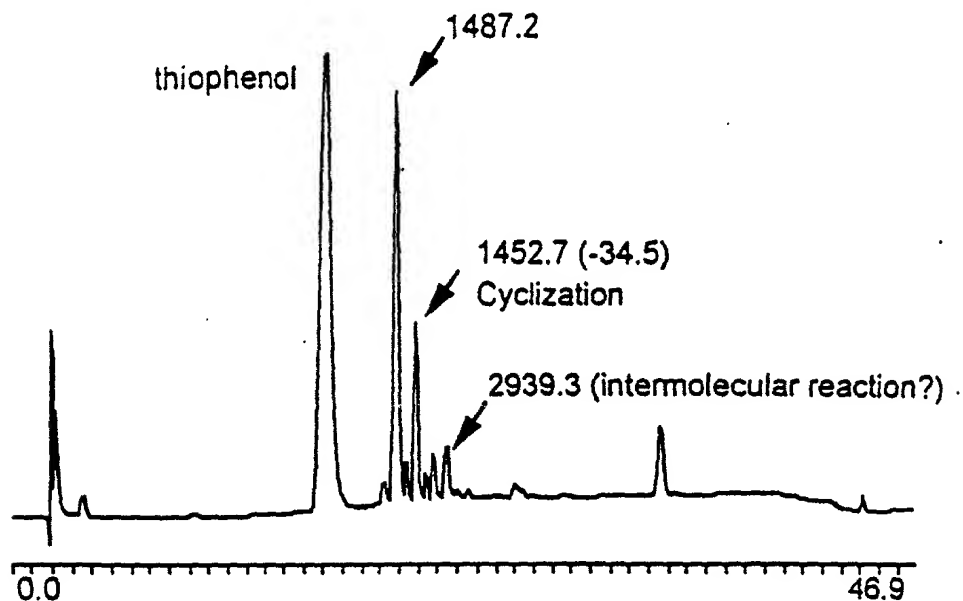


FIG. 2A

2. In the presence of a thioester peptide

H-CGFRVREFGDNTA-COSH MW=1487.6 + *H*-DSVISLSGDH-SPAL MW= 1230.2

MW of Ligation product = 2498.7

6M GU·HCL, 0.1M NaPi, 0.5% thiophenol, room temperature, overnight

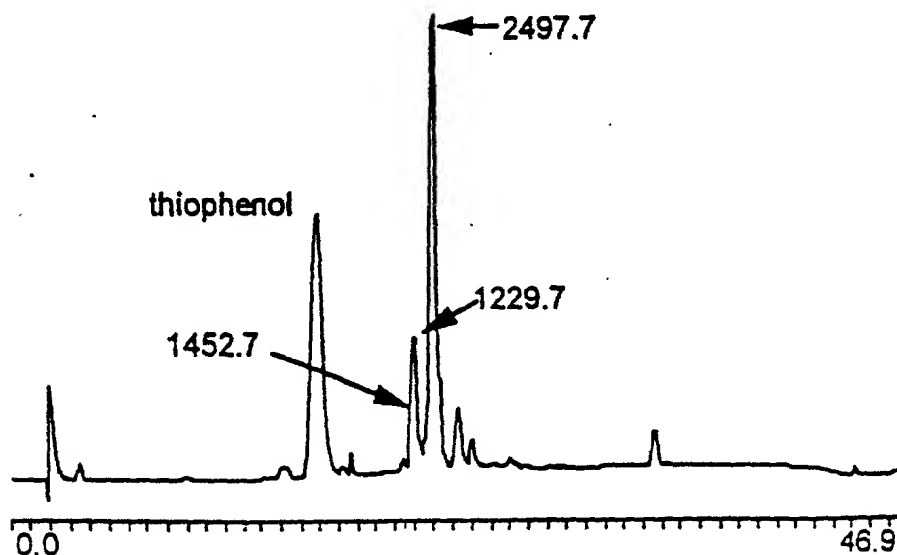


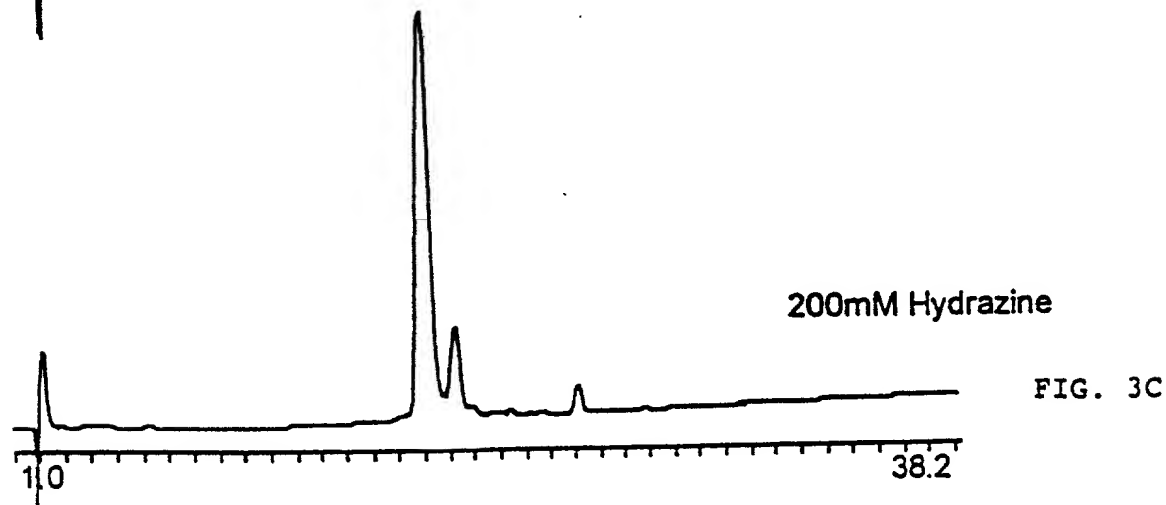
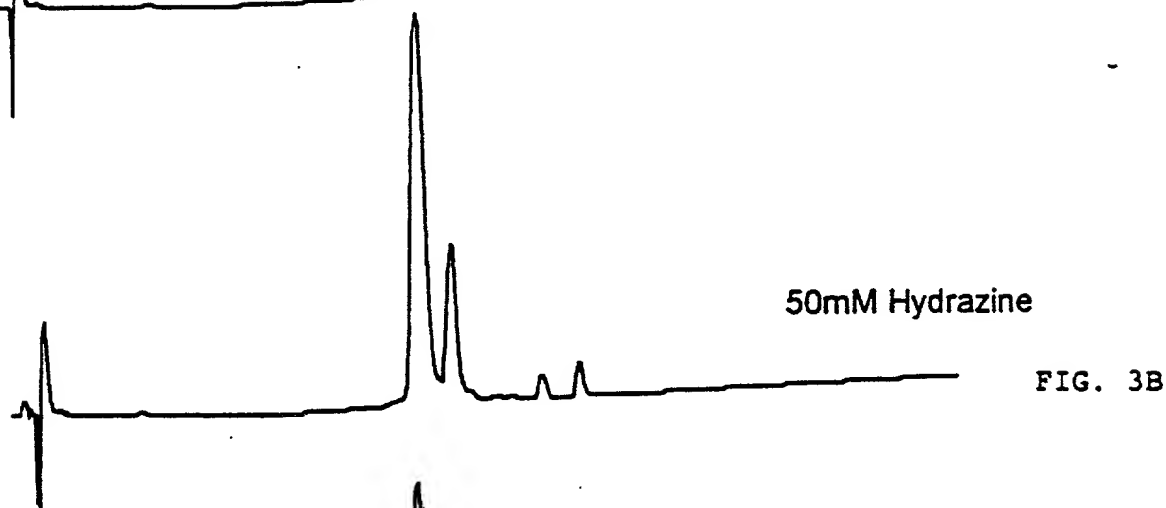
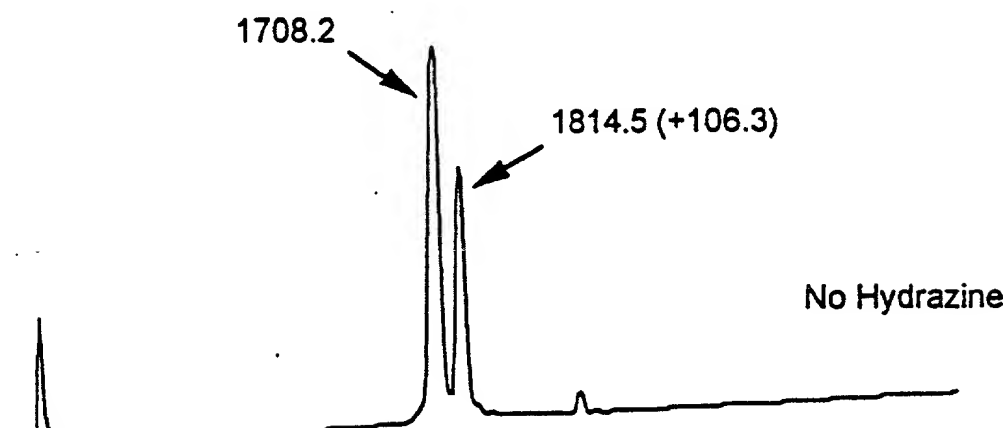
FIG. 2B

MSC Removal Experiments

MSC-CTSAGPHFNPLSRKHG-OH MW=1859.1

H-CTSAGPHFNPLSRKHG-OH MW=1708.9

Aliquot of peptide in 6M Gu·HCl, 0.1M
NaPi, pH 7.5 was diluted into 1N NaOH for
two minutes, quenched with 1N HCl



MSC Removal Experiments (Cont'd)

Lev-MSC-LTEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHG-COSH

MW=4022.4

H-LTEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHG-COSH

MW=3745.1

Aliquot of peptide in 6M Gu•HCl, 0.1M NaAc, pH 4.6 was diluted into 6M Gu•HCl, 0.1M NaAc, pH 14 for two minutes, quenched with 6M Gu•HCl, 0.1M NaAc, pH 2.0

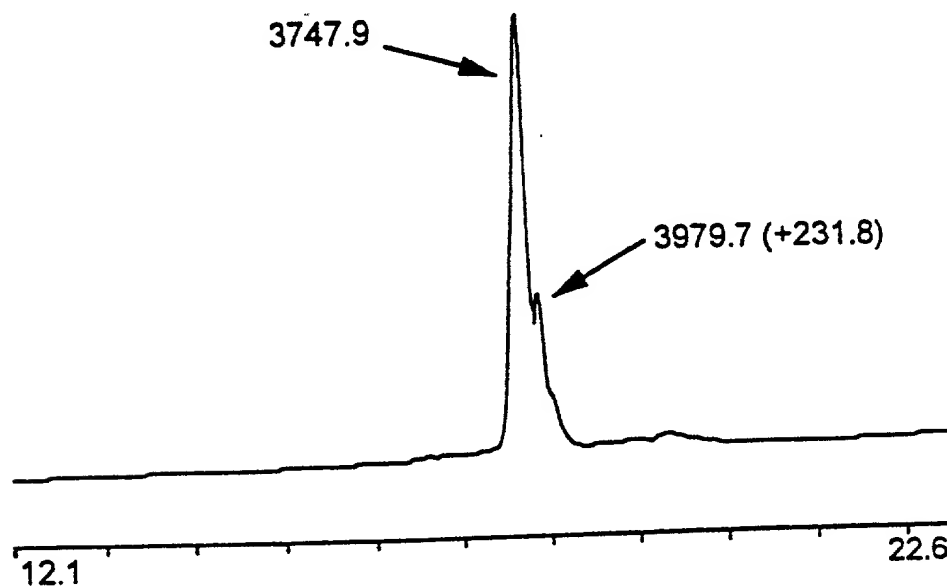


FIG. 4

Resin Preparation

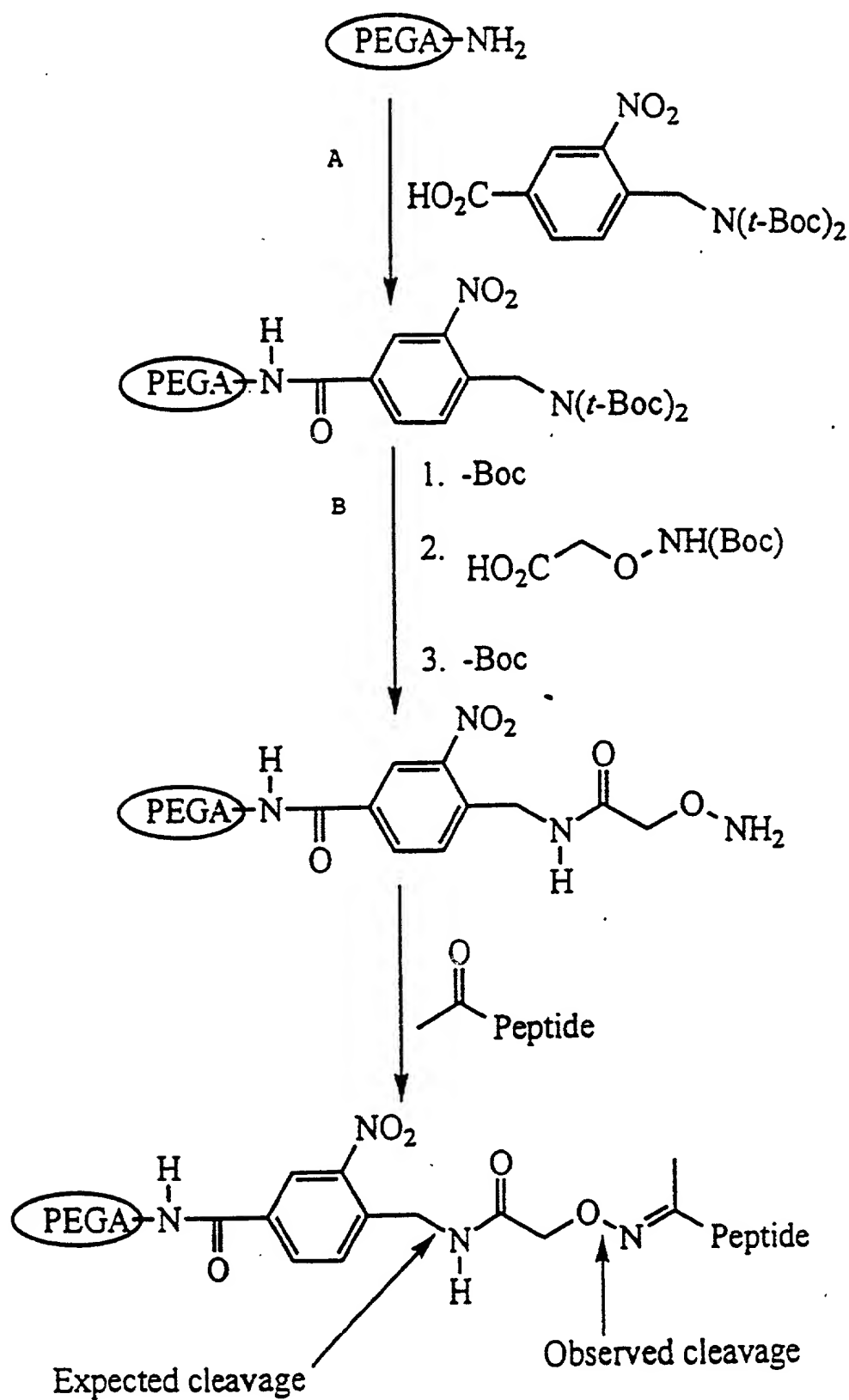
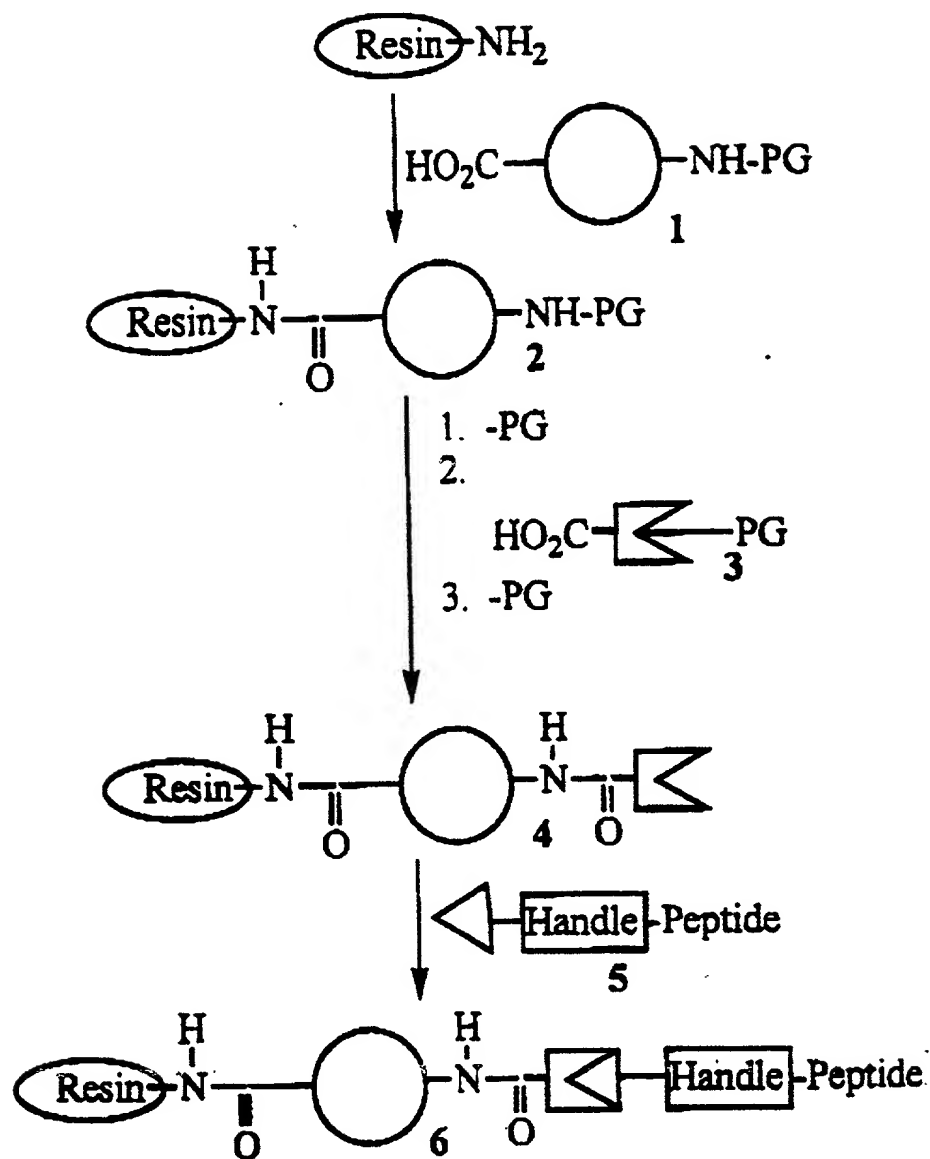


FIG. 5A

Resin Preparation



$\text{HO}_2\text{C}-\text{Circle}-\text{NH}-$ = cleavable linker used for monitoring with Maldi, Electrospray Mass Spec, etc...

PG = protecting group

$\text{HO}_2\text{C}-\text{Triangle}-$ = functional group added to resin to couple with peptide

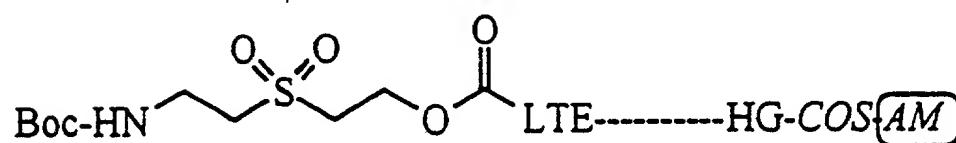
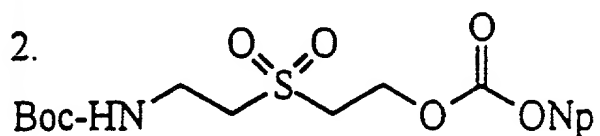
$\text{Triangle}-\text{Handle}-\text{Peptide}$ = peptide functionalized with 1) a cleavable handle for release of peptide/protein from the resin at completion of synthesis and 2) functional group to couple to resin

FIG. 5B

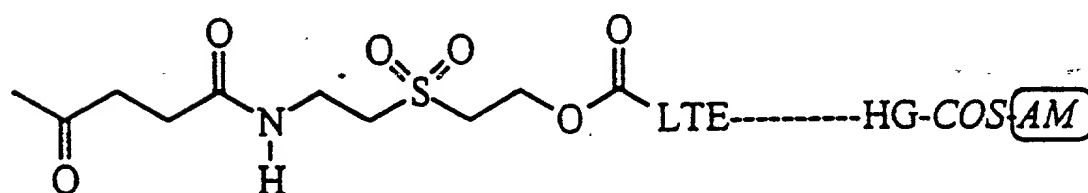
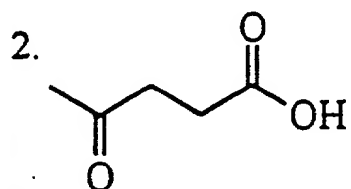
**Derivatization of Segment 1
(N-terminal)**



1. -Boc



1. -Boc



HF Cleavage

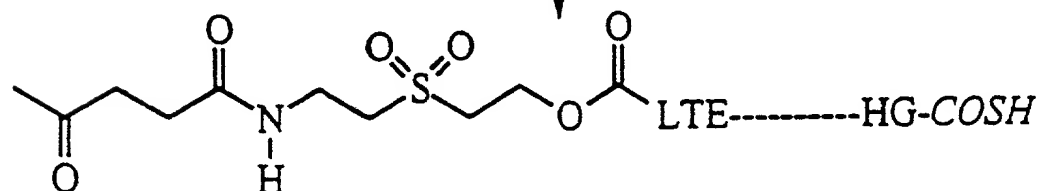


FIG. 6

Polymer-Supported Ligation on PEGA

Lev-*MSC-LTEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHG-COSH* (1)
+ Resin-PCL-ONH₂

↓ 1. pH 4.6, 6M Gu·HCl, 0.1 acetate

Resin-PCL-oxime-MSC-LTEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHG-COSH (1)

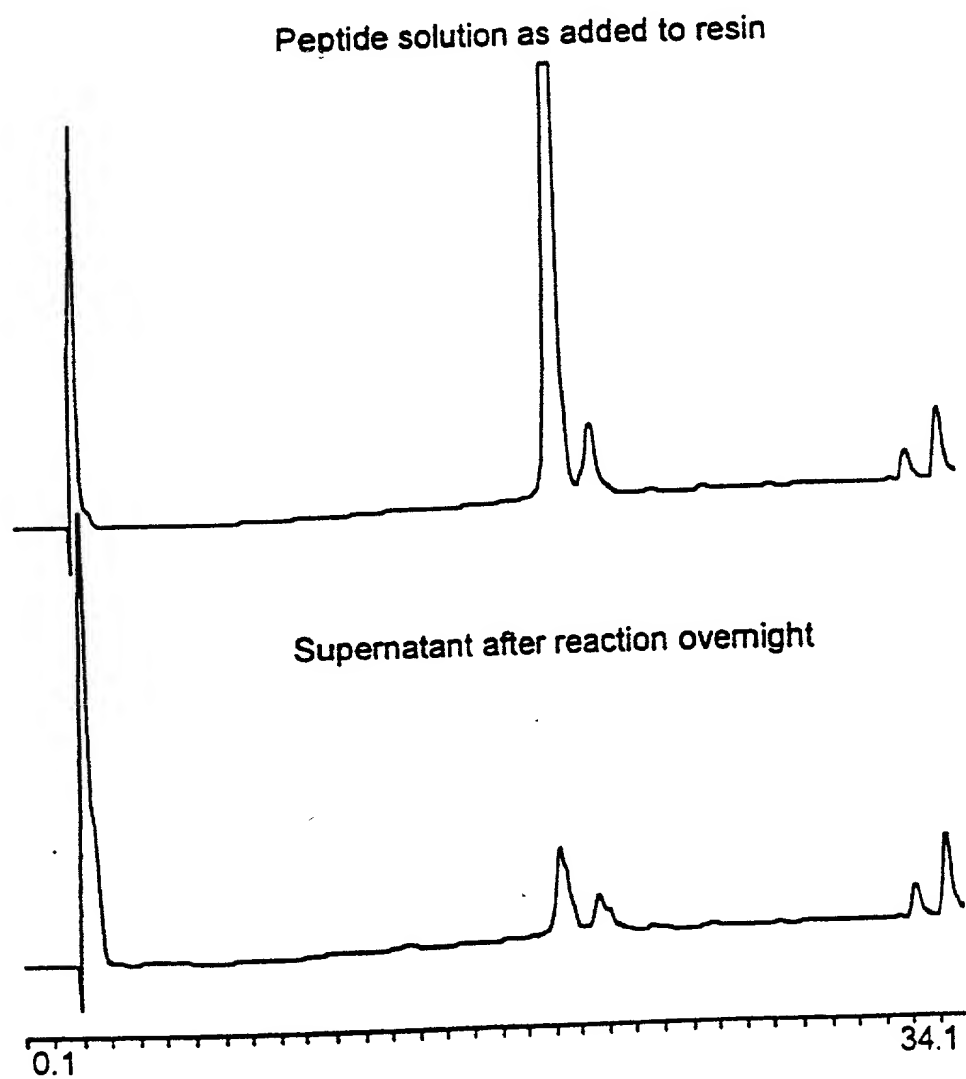


FIG. 7A

FIG. 7B

Polymer-Supported Ligation on *ISeo*

Lev-*MSC*-LTEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHG-COSH(1)
+ Resin-PCL-ONH₂

↓ 1. pH 4.6, 6M Gu·HCl, 0.1 acetate

Resin-PCL-oxime-*MSC*-LTEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHG-COSH(1)

Peptide solution as added to resin



FIG. 8A

Supernatant after reaction overnight

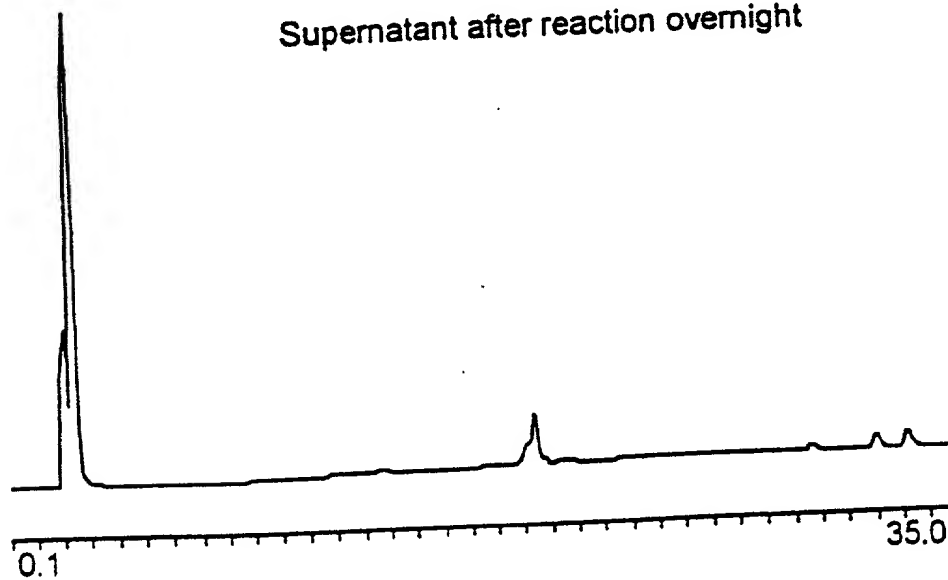


FIG. 8B

Polymer-Supported Ligation on Isco

Lev-MSC-LTEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHG-COSH(1)
+ Resin-PCL-ONH₂

↓ 1. pH 4.6, 6M Gu·HCl, 0.1 acetate

Resin-PCL-oxime-MSC-LTEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHG-COSH(1)
Maldi Mass = 4022, Base Cleavage Mass = 3745

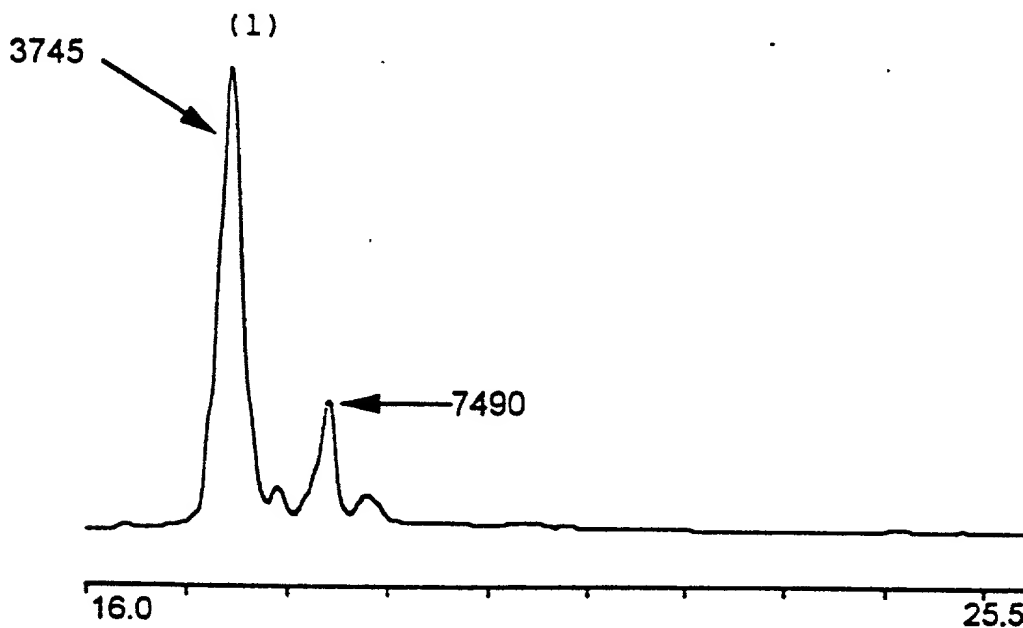


FIG. 9A

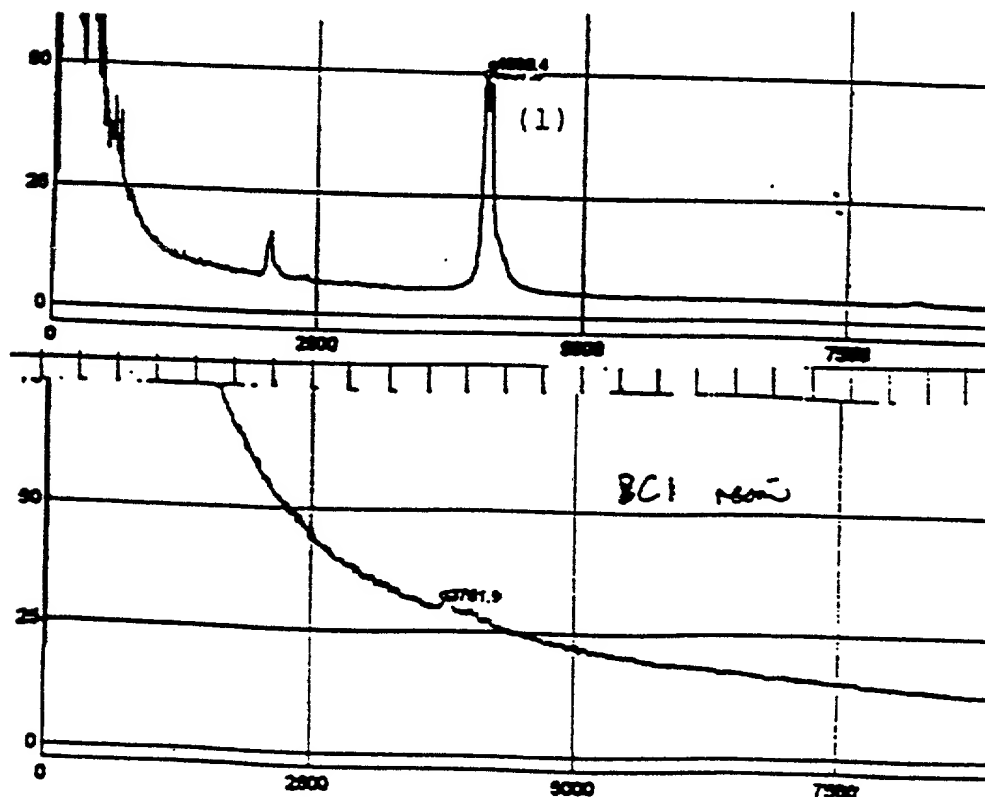


FIG. 9B

FIG. 9C

Polymer-Supported Ligation *ONT* sco

Resin-PCL-oxime-*MSC*-L-*TEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHG-COSAc* (1)

Maldi Mass = 4080, Base Cleavage Mass = 3729

+ *H-CGFRVREFGDNTA-COSH* (2)

↓ 3. pH 7.5, 6M Gu•HCl, 0.1M phosphate, 0.5% thiophenol

Resin-PCL-oxime-*MSC*-L-*TEGLHGFHVHEFGDNTAGCTSAGPHFNPLSRKHGCGFRVREF-GDNTA-COSH* (1+2)

Maldi Mass = 5476, Base Cleavage Mass = 5199

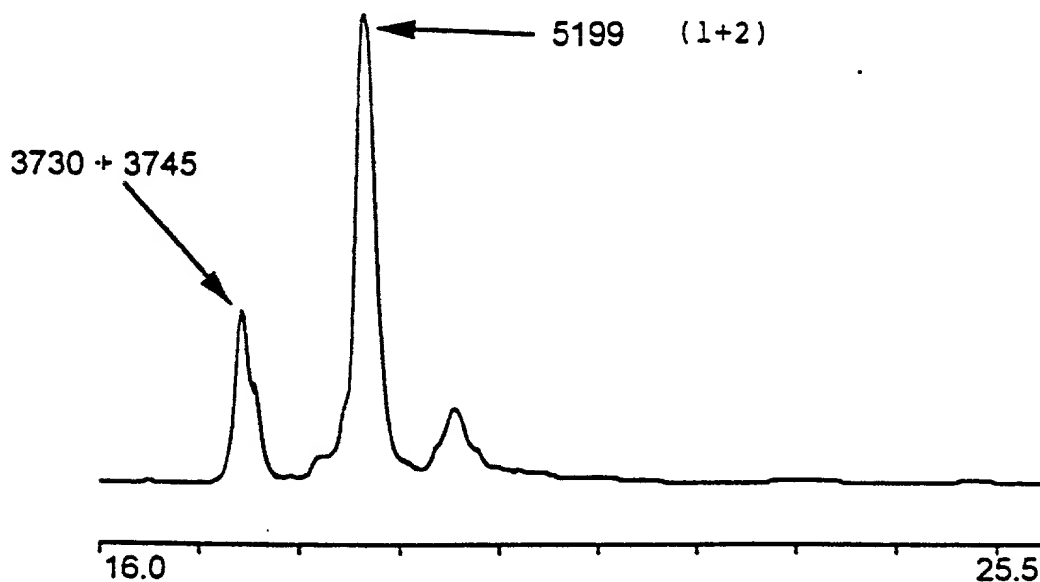


FIG. 10
A

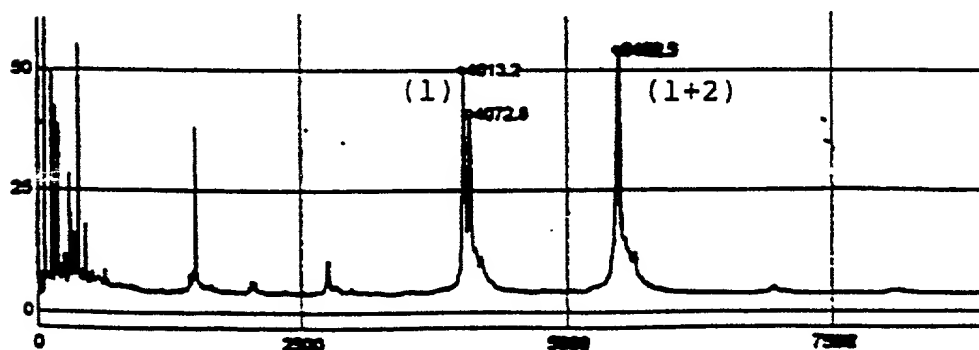


FIG. 10
B

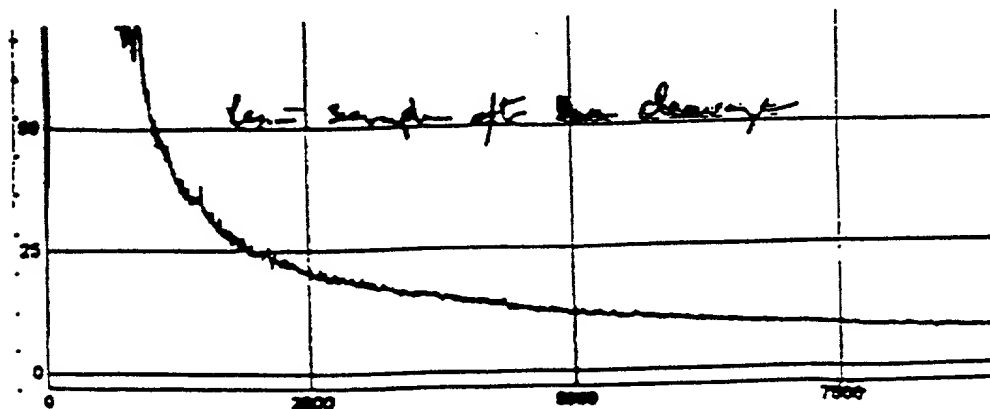


FIG. 10
C

Polymer-Supported Ligation on Isco

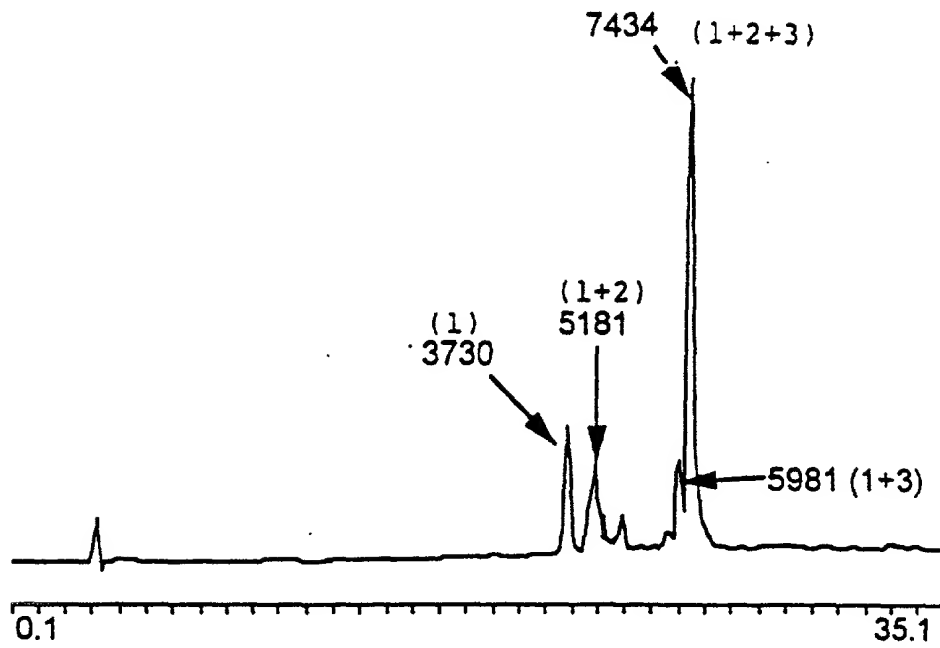


FIG. 11

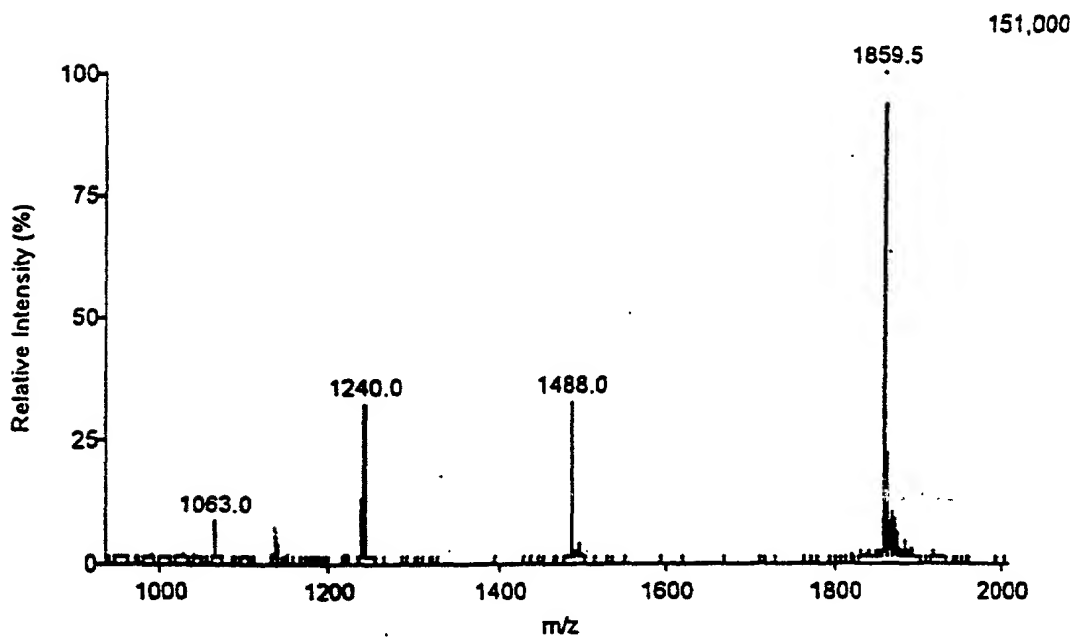


FIG. 12A

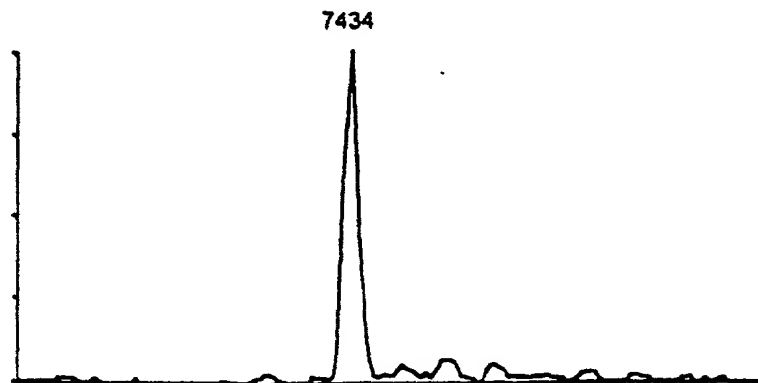


FIG. 12B

Polymer-Supported Ligation on PEG6A

No photocleavable linker

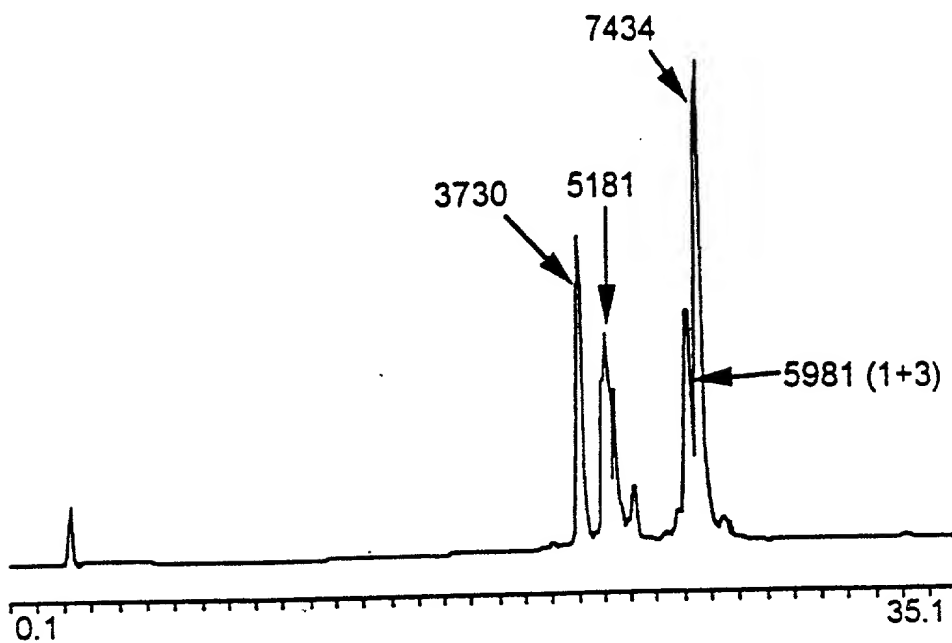


FIG. 13

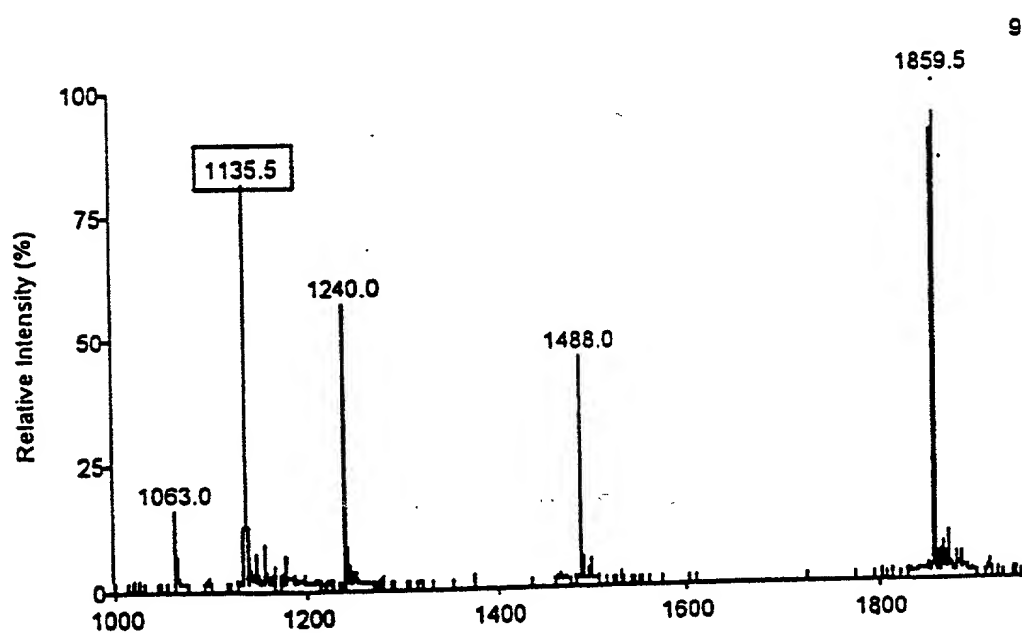


FIG. 14A

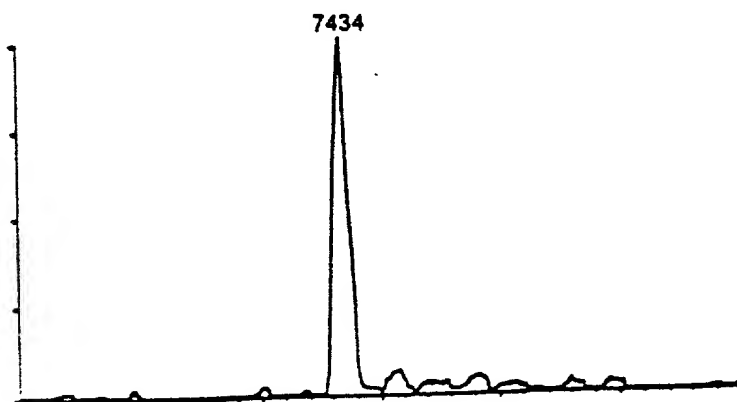


FIG. 14B

On Resin Purification

Solution processing

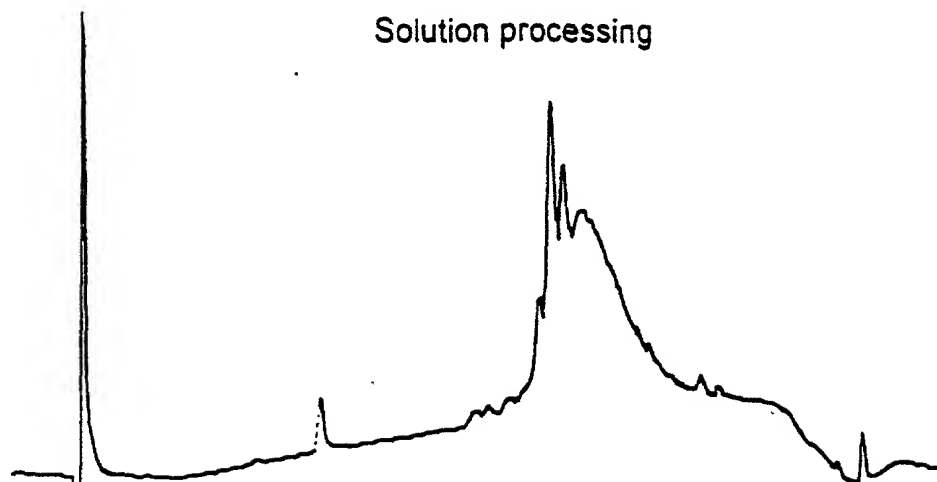


FIG. 15A

DNP removal

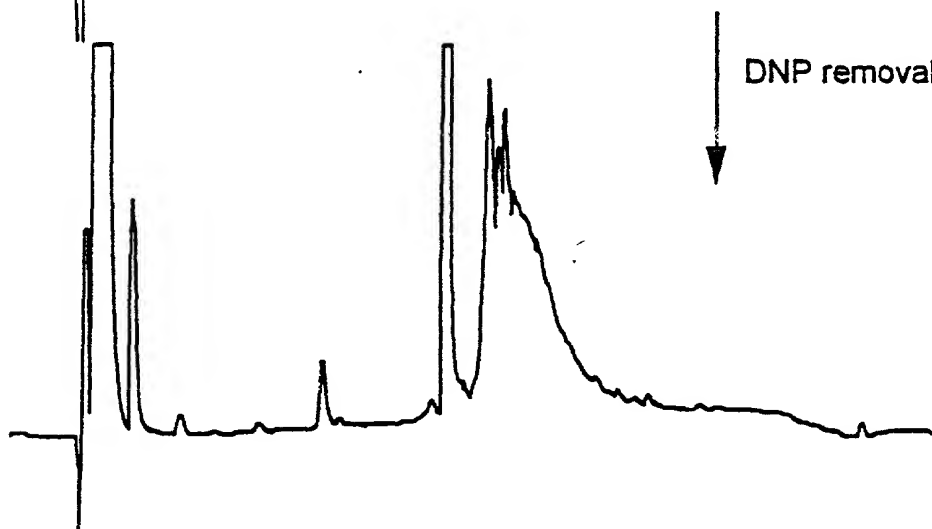


FIG. 15B

0.0 46.8

Polymer-supported processing

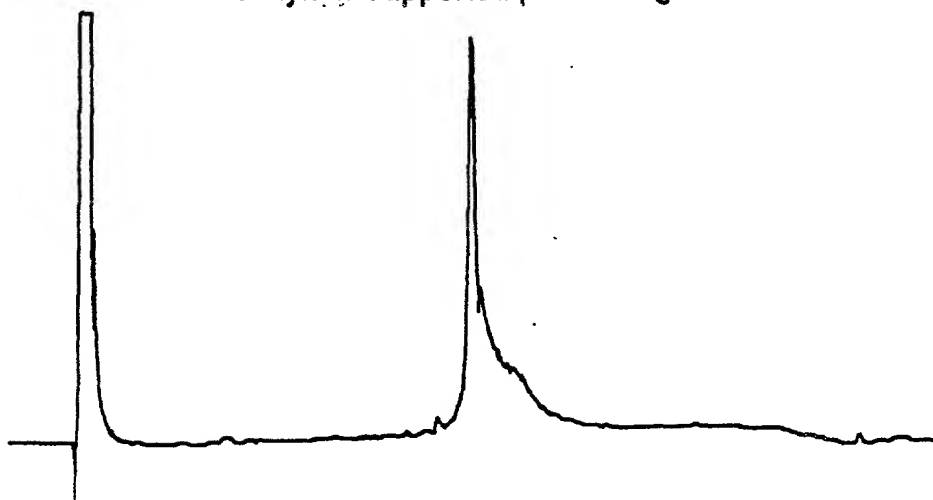


FIG. 15C

0.1 46.8

FIG. 15A

Synthesis of MIF by Solid Phase Native Ligations

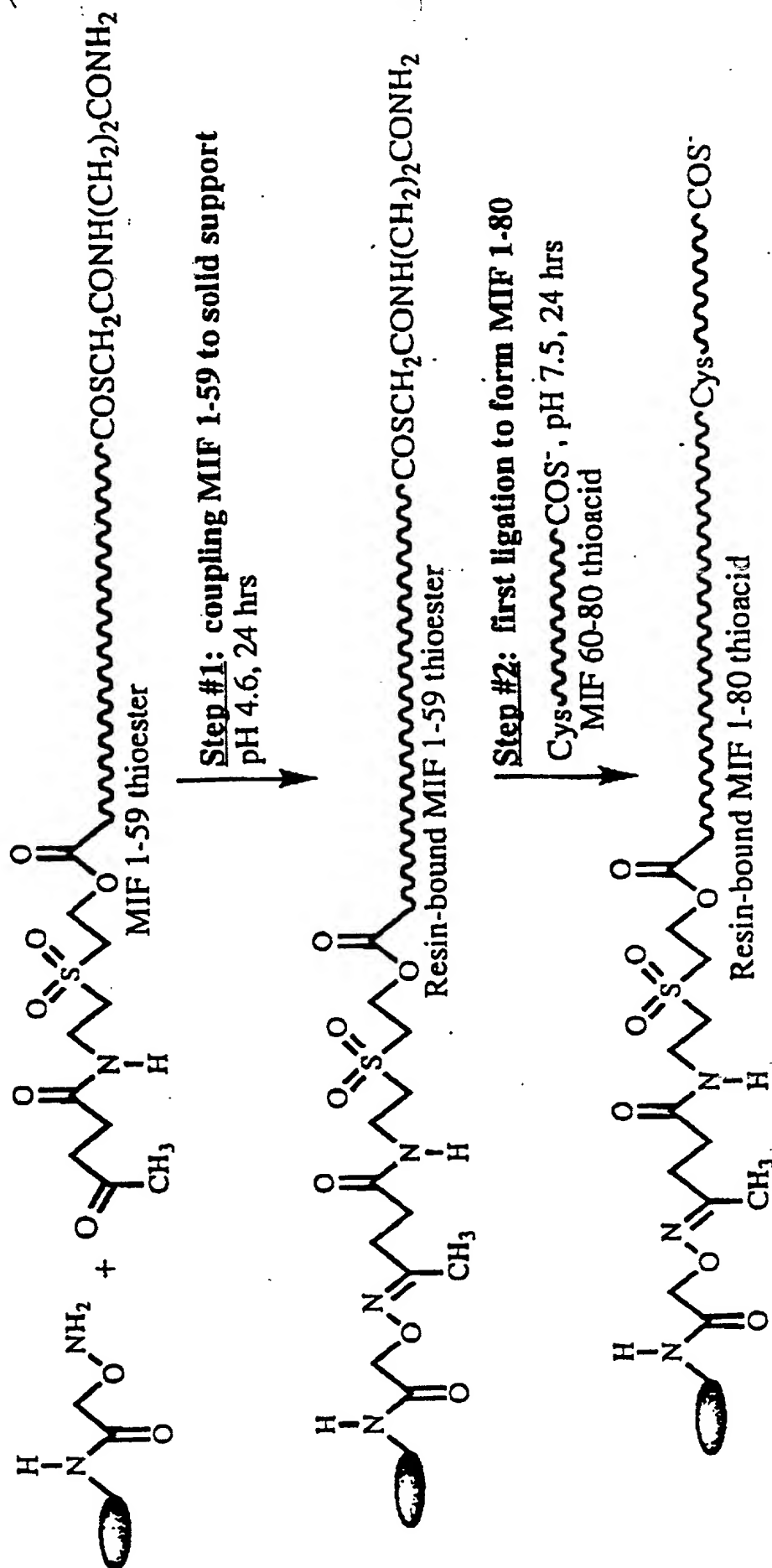


FIG. 16A

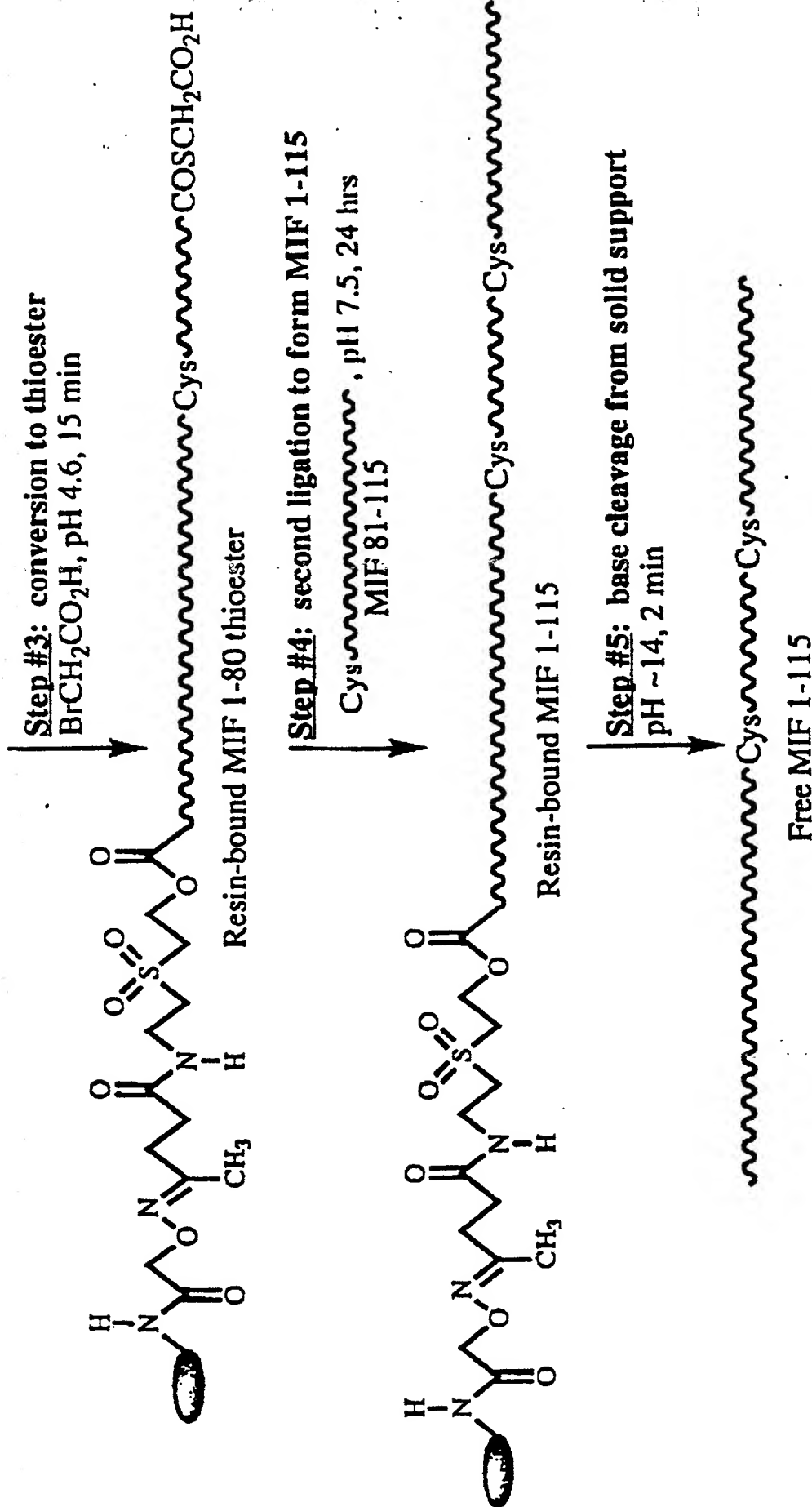


FIG. 16B

Modification of N-terminal Peptide Segment and Solid Support

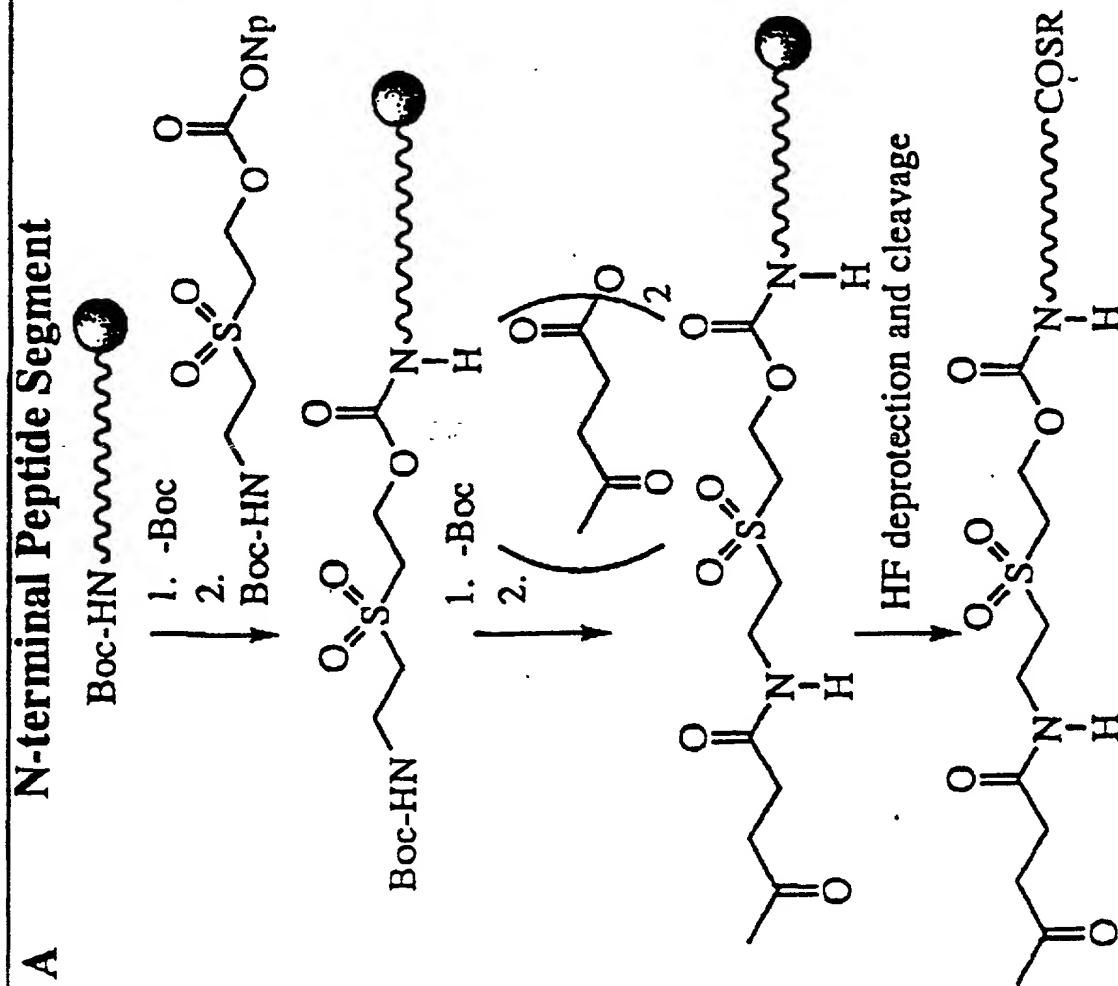


FIG 17A

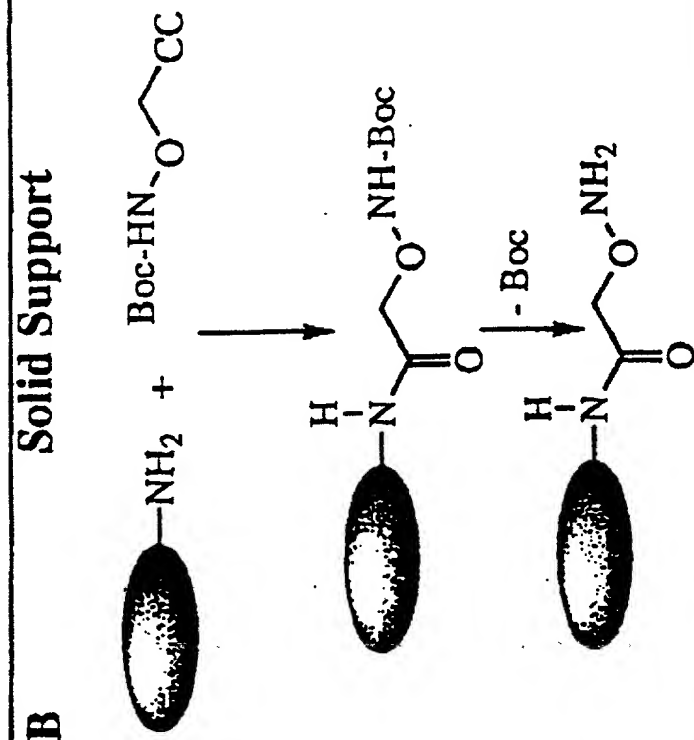
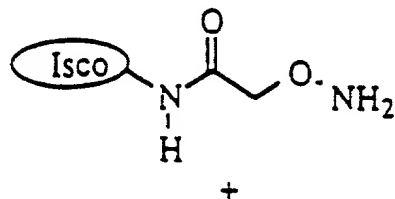


FIG. 17B

Coupling of MIF 1-59 to Solid Support

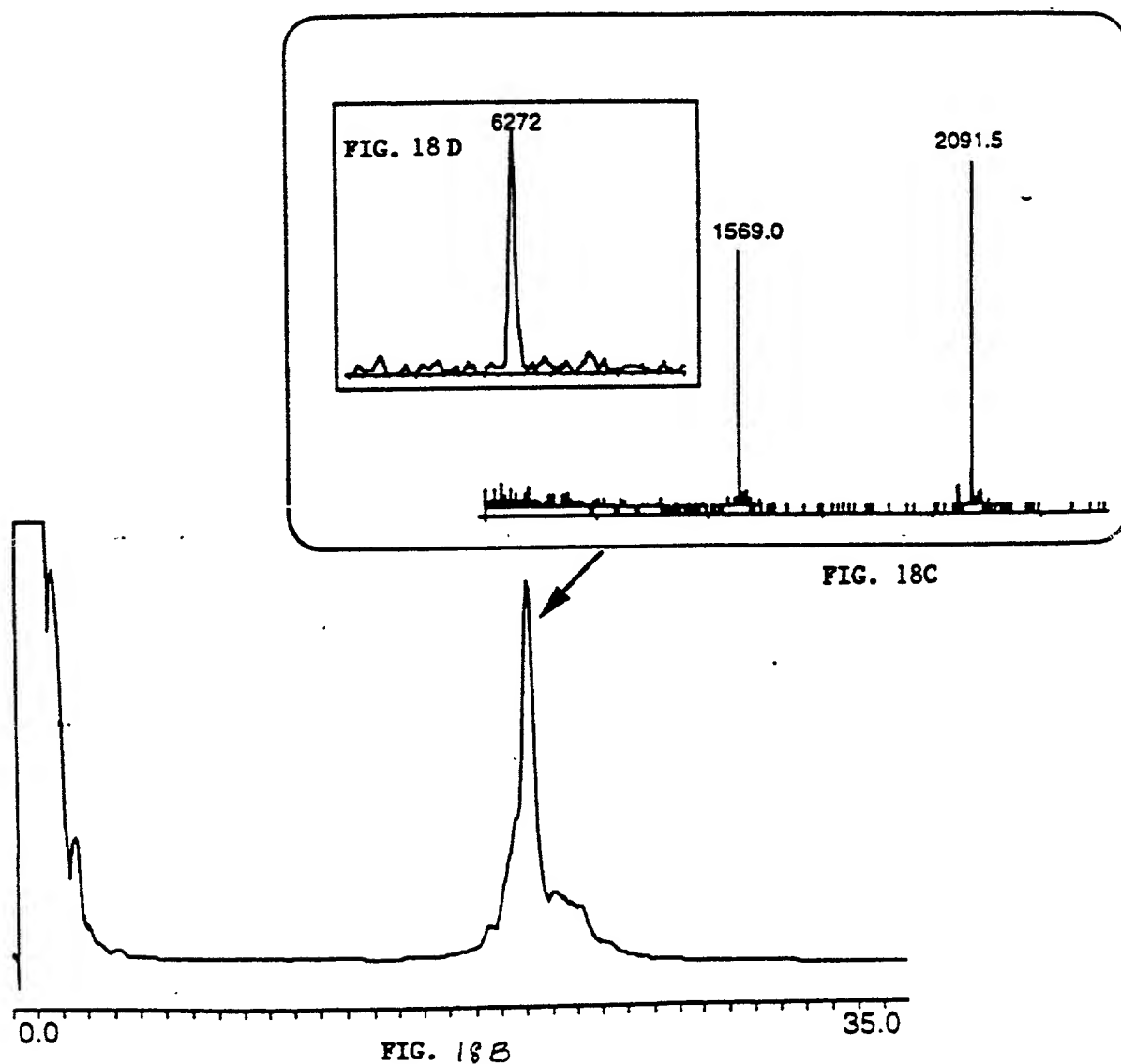


Ketone-MS handle-Met¹-MIF 2-58-Leu⁵⁹-SAc-βAla-CO₂H

#1

Isco - Oxime-MS handle-Met¹-MIF 2-58-Leu⁵⁹-SAc-βAla-CO₂H
 Expected base cleavage mass = 6271

FIG. 18A



Ligation to form MIF¹⁻⁸⁰

Isco — Oxime-MS handle-Met¹-MIF 2-58-Leu⁵⁹-SAc-βAla-CO₂H

#2
↓
Cys⁶⁰-MIF 61-79-Leu⁸⁰-COSH

Isco — Oxime-MS handle-Met¹-MIF 2-79-Leu⁸⁰-COSH
Expected base cleavage mass = 8502

FIG. 19A

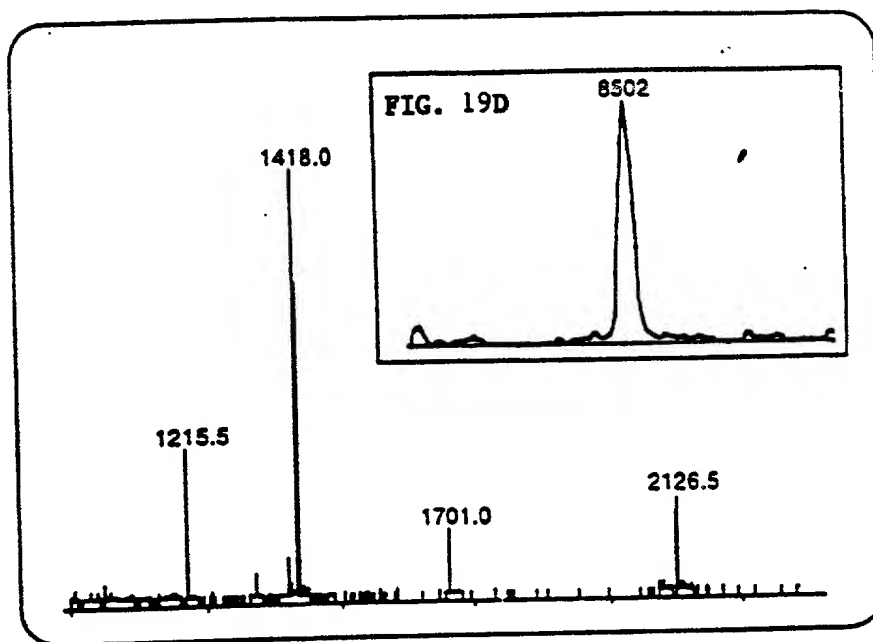


FIG. 19C

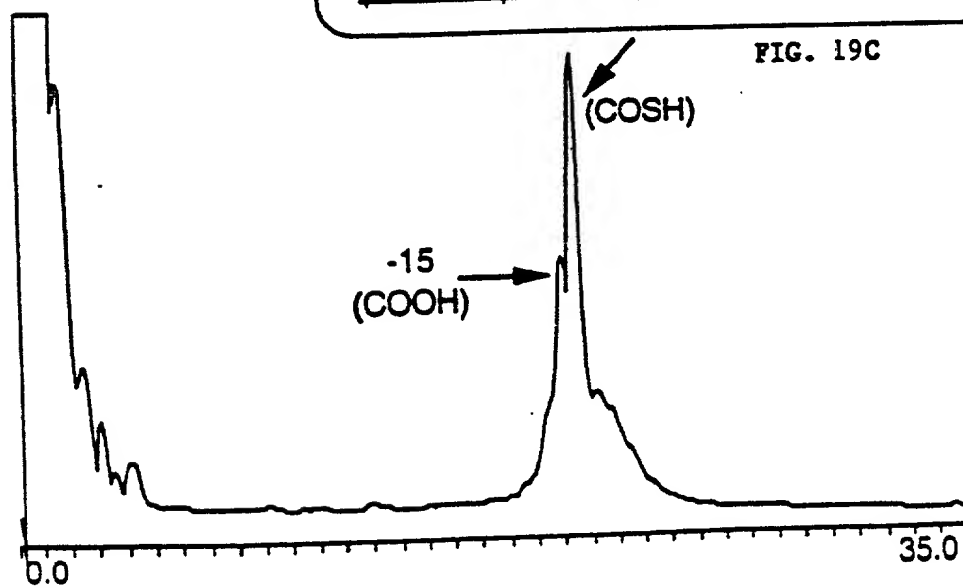


FIG. 19B

(Ligation to form MIF 1-115

Isco — Oxime-MS handle-Met¹-MIF 2-79-Leu³⁰-COSAc

#4
Cys⁸¹-MIF 82-114-Ala¹¹⁵-CO₂H
6M Gu-HCl, 0.1 M Na Pi, 0.5% thiophenol
0.15 M Methionine, pH 7.5

Isco — Oxime-MS handle-Met¹-MIF 2-114-Ala¹¹⁵-CO₂H
Expected base cleavage mass = 12450

FIG. 20A

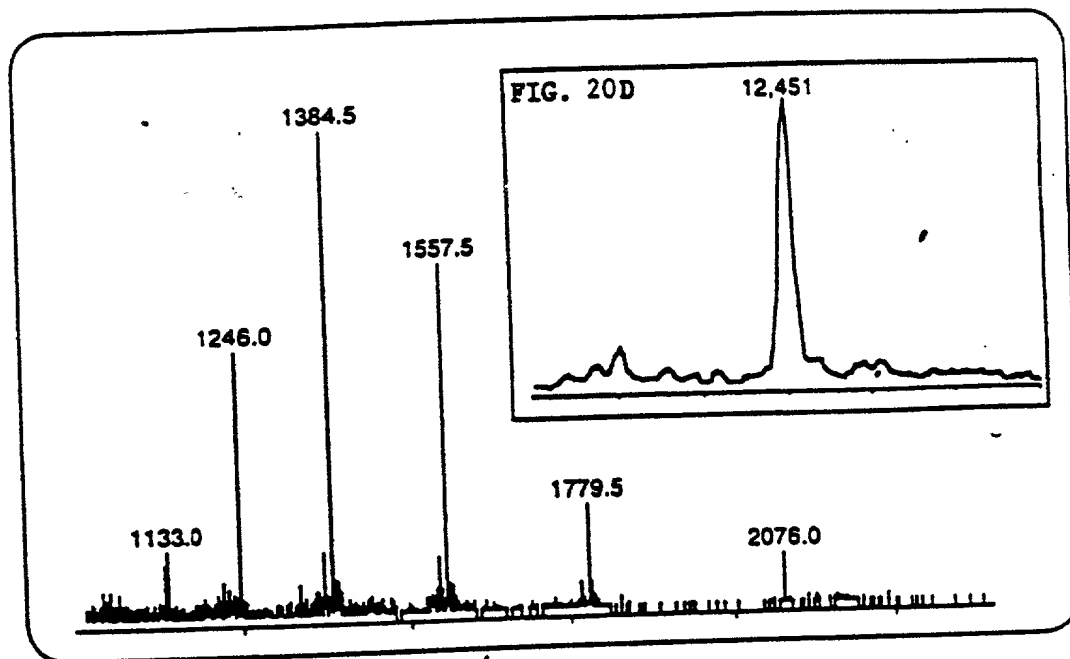


FIG. 20C

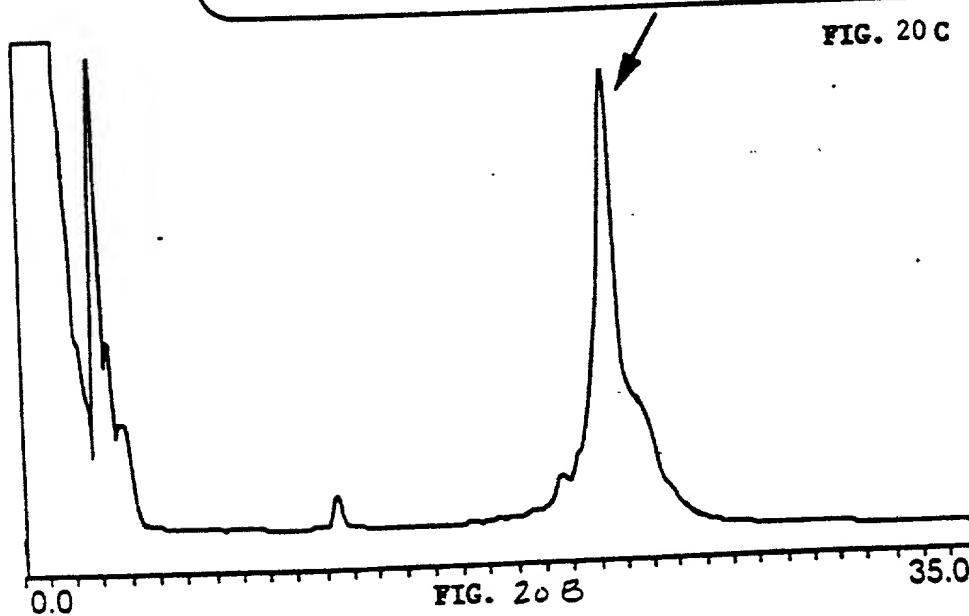


FIG. 20B

Solid Phase Chemical Ligations in the C- to N-terminal Direction

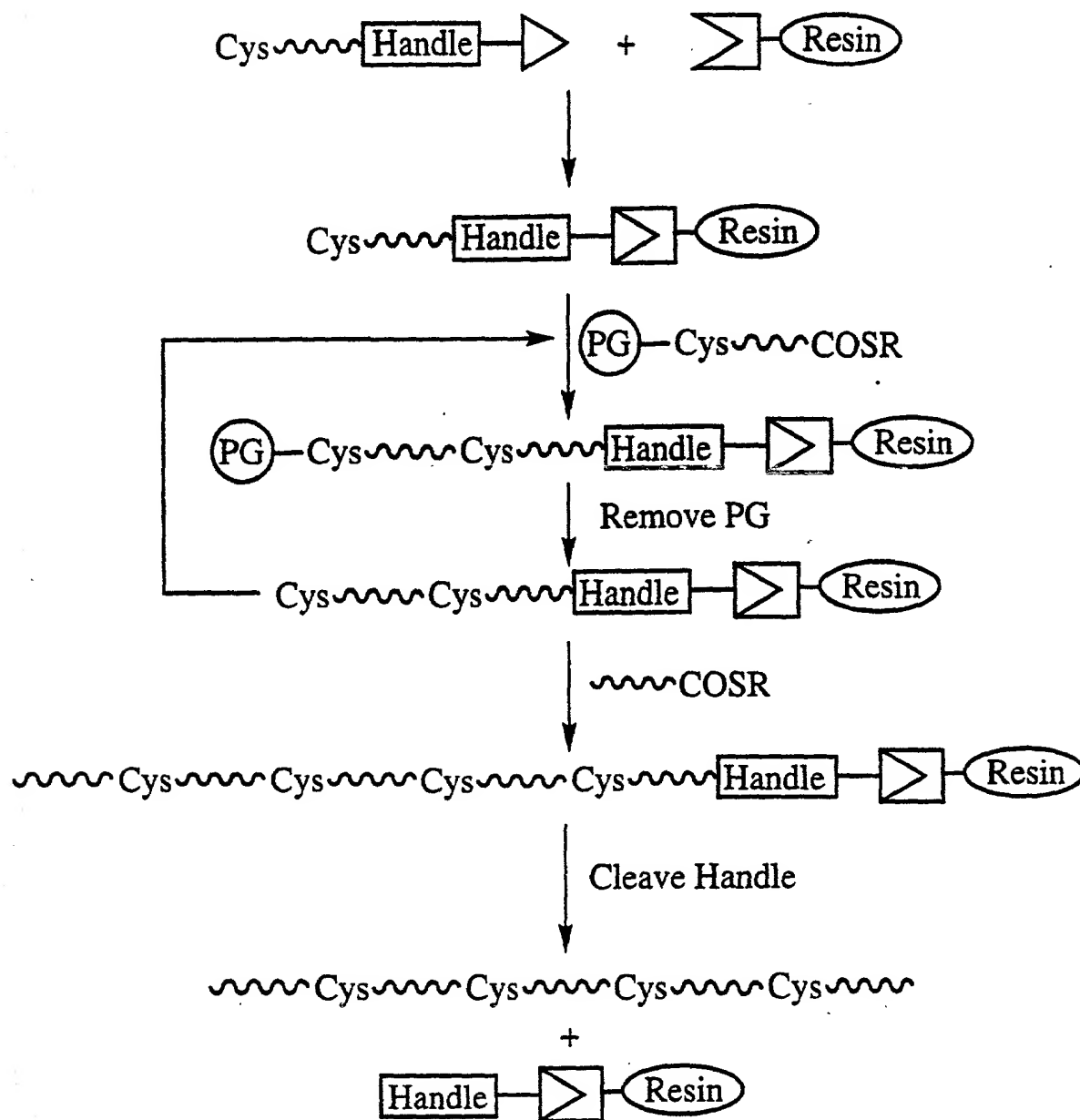


FIG 21

Solid Phase Chemical Ligations in the C- to N-Terminal Direction **Synthesis of Phospholipase A2, Group 5 (PLA2G5)**

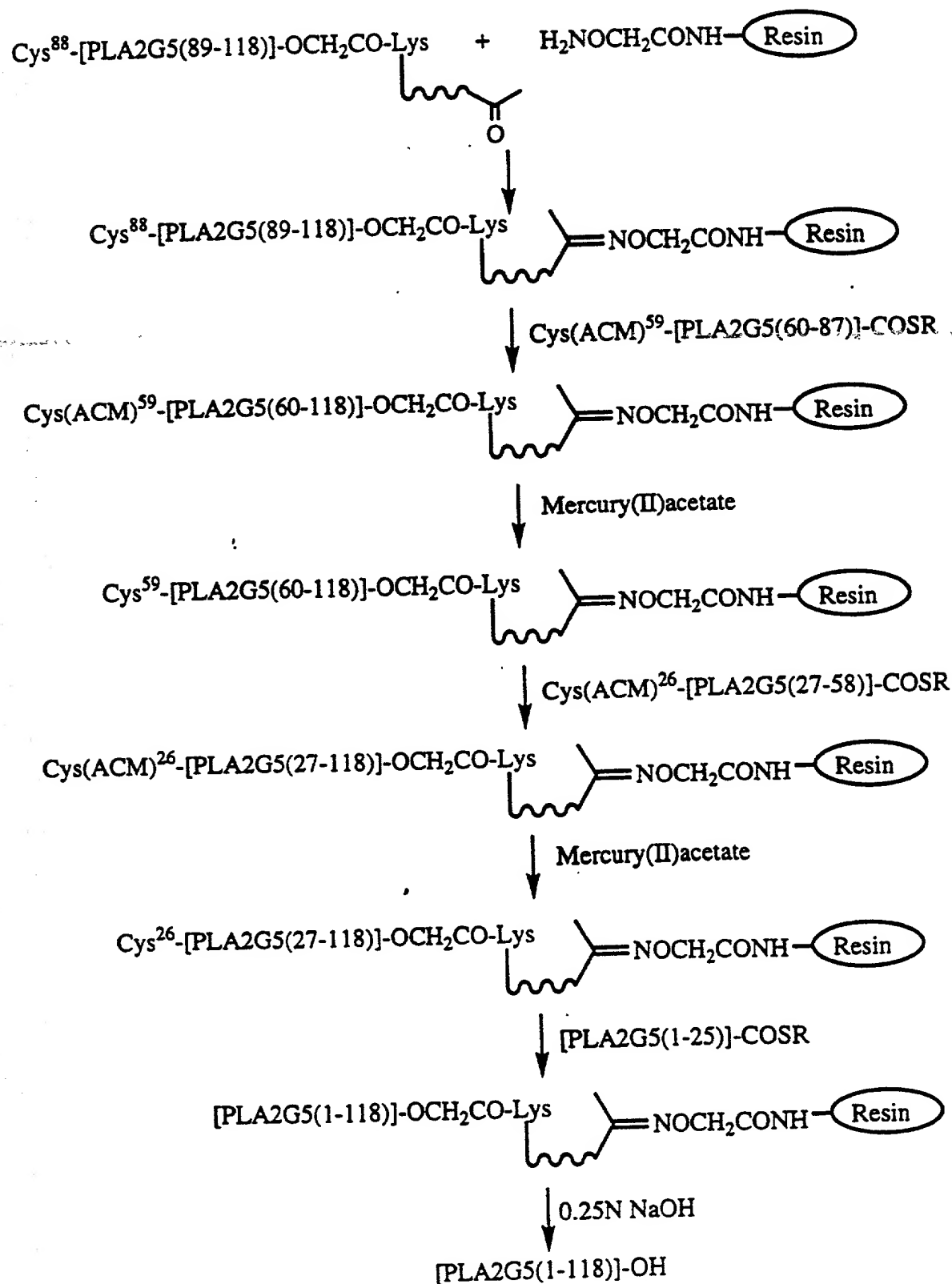


Figure 22

Synthesis of Cam ester derivative

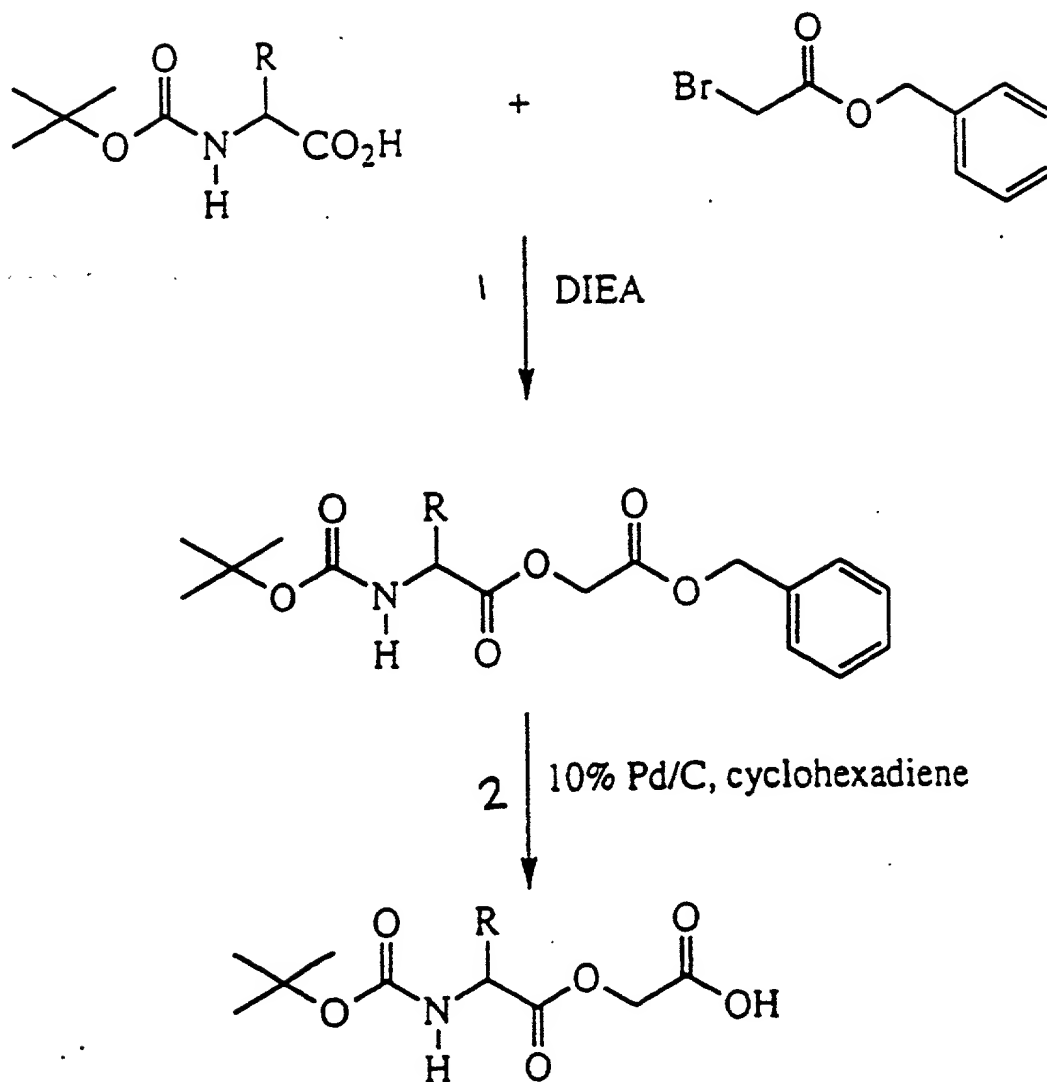


FIG. 23

Synthesis of C-terminal peptide segment

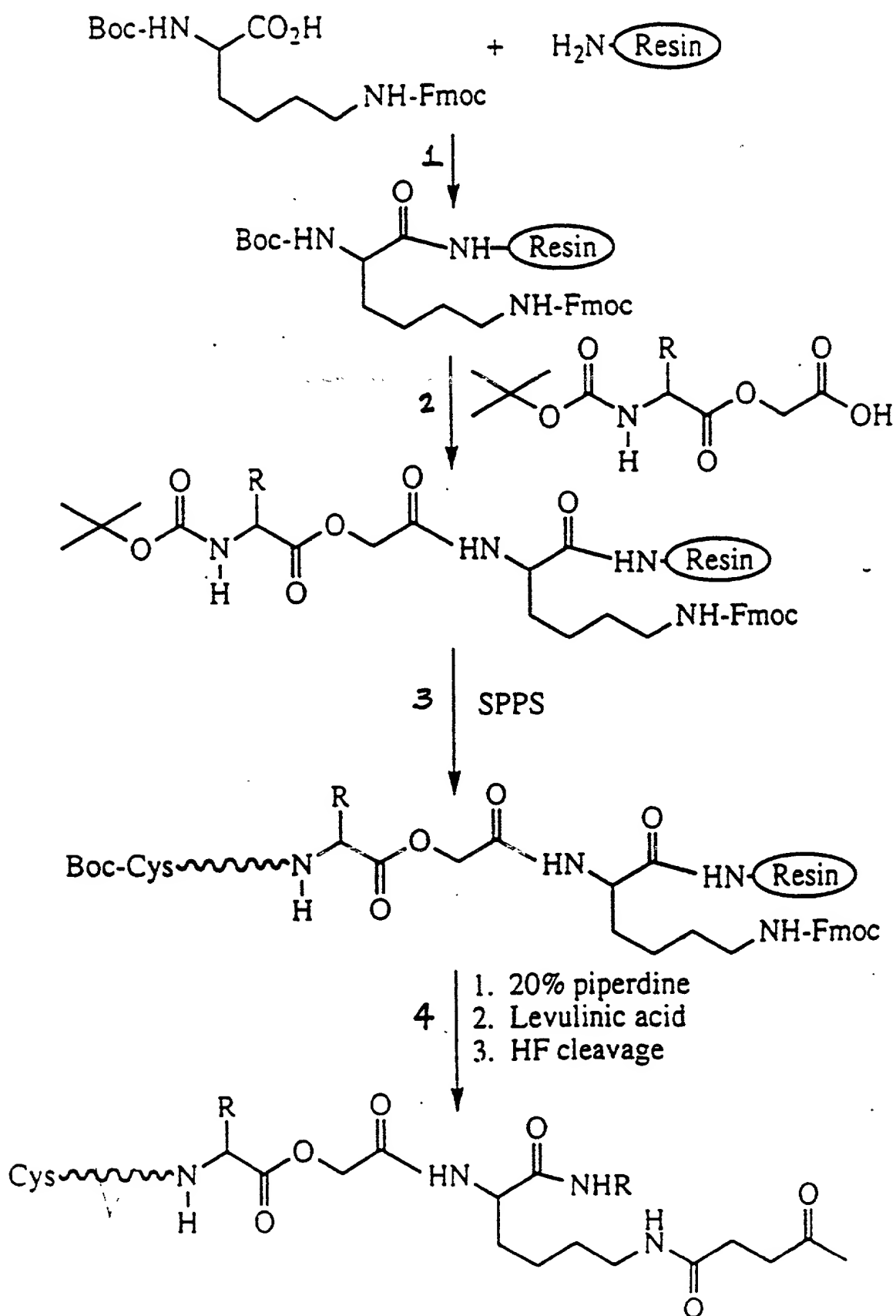


FIG. 24

Universal Solid Phase Chemical Ligation
(Bidirectional Ligations: C- to N-Terminal Ligations First)

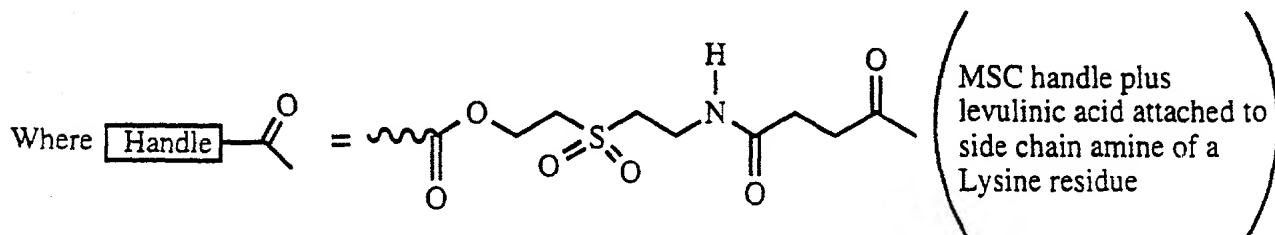
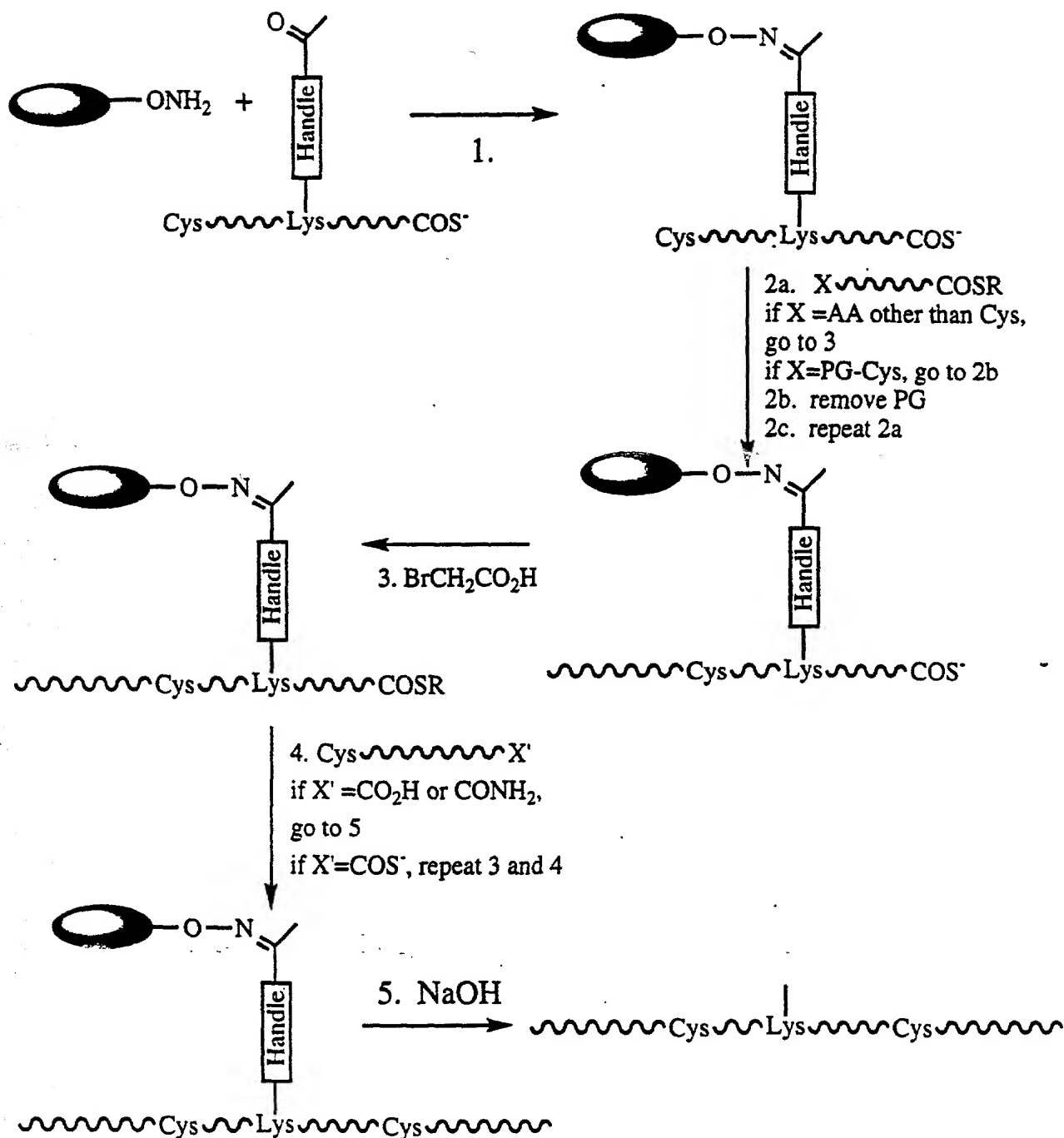


FIG. 25A

Universal Solid Phase Chemical Ligation **(Bidirectional Ligations: N- to C-Terminal Ligations First)**

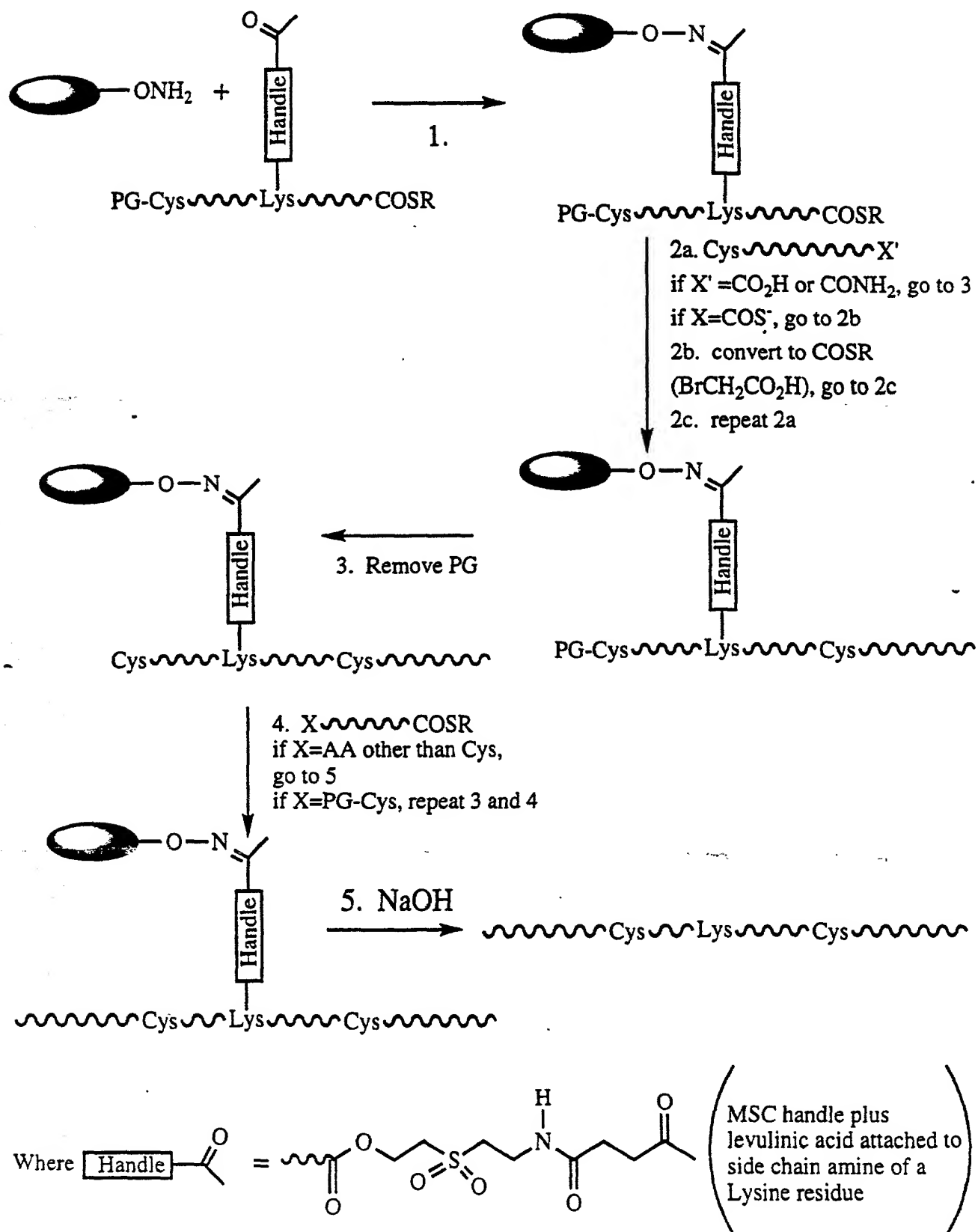


FIG. 25B

Synthesis of Modified Peptide Segment for Universal Solid Phase Chemical Ligation

Starting with an appropriate resin (thioester or thioacid generating), synthesize the peptide using standard Boc protocols until the Lys residue of choice is reached. Couple a Boc-Lys(Fmoc)-OH, then continue the synthesis.

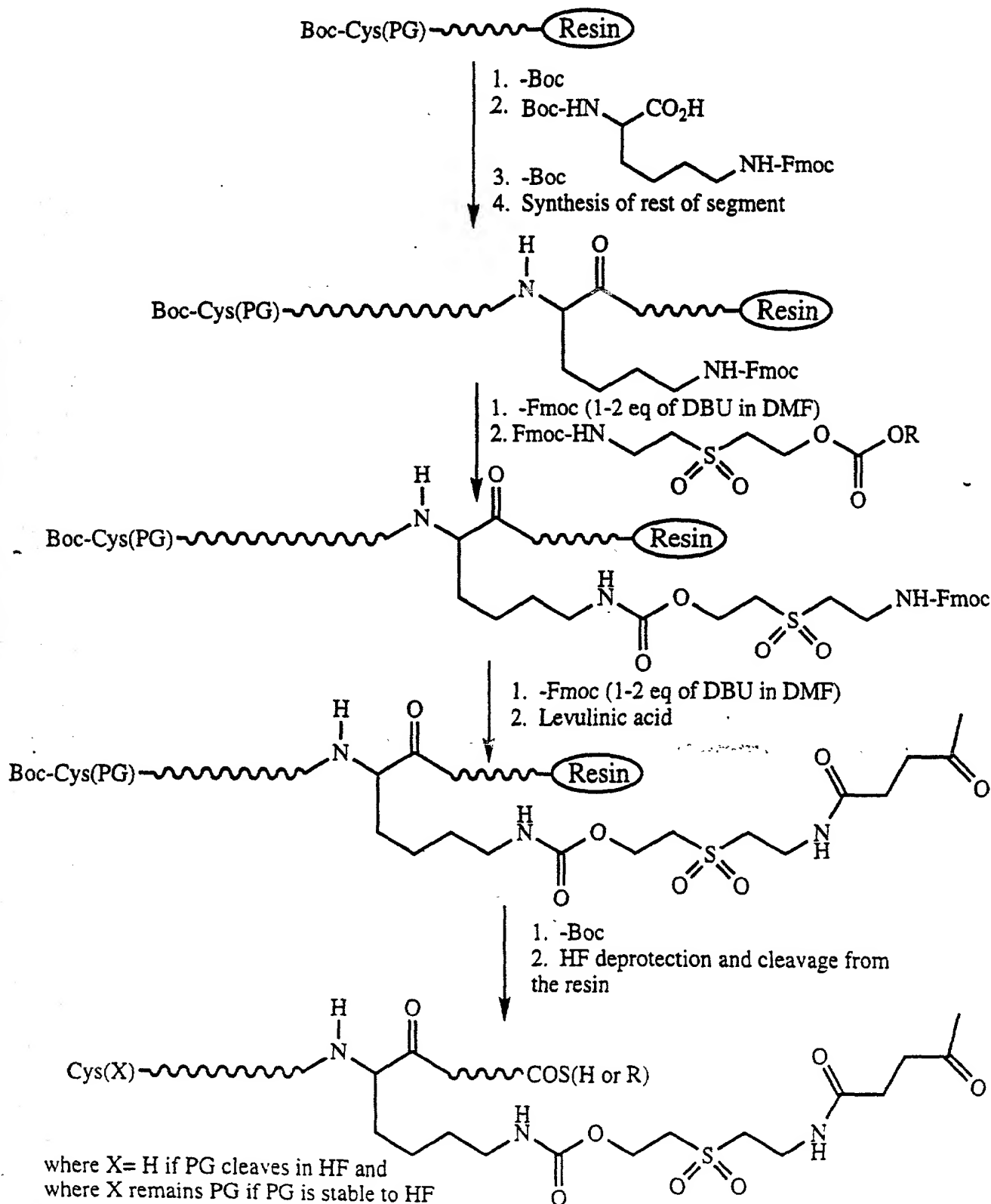


FIG. 25C

1 21 47
TLQKKIEEIAAKYKHSVVKKCCYDGACVNNDETCEQRAARISLGPKCIKAFTECC
VVASQLRANISHKDMQLGR
74



Synthesis of C-terminal peptide segment

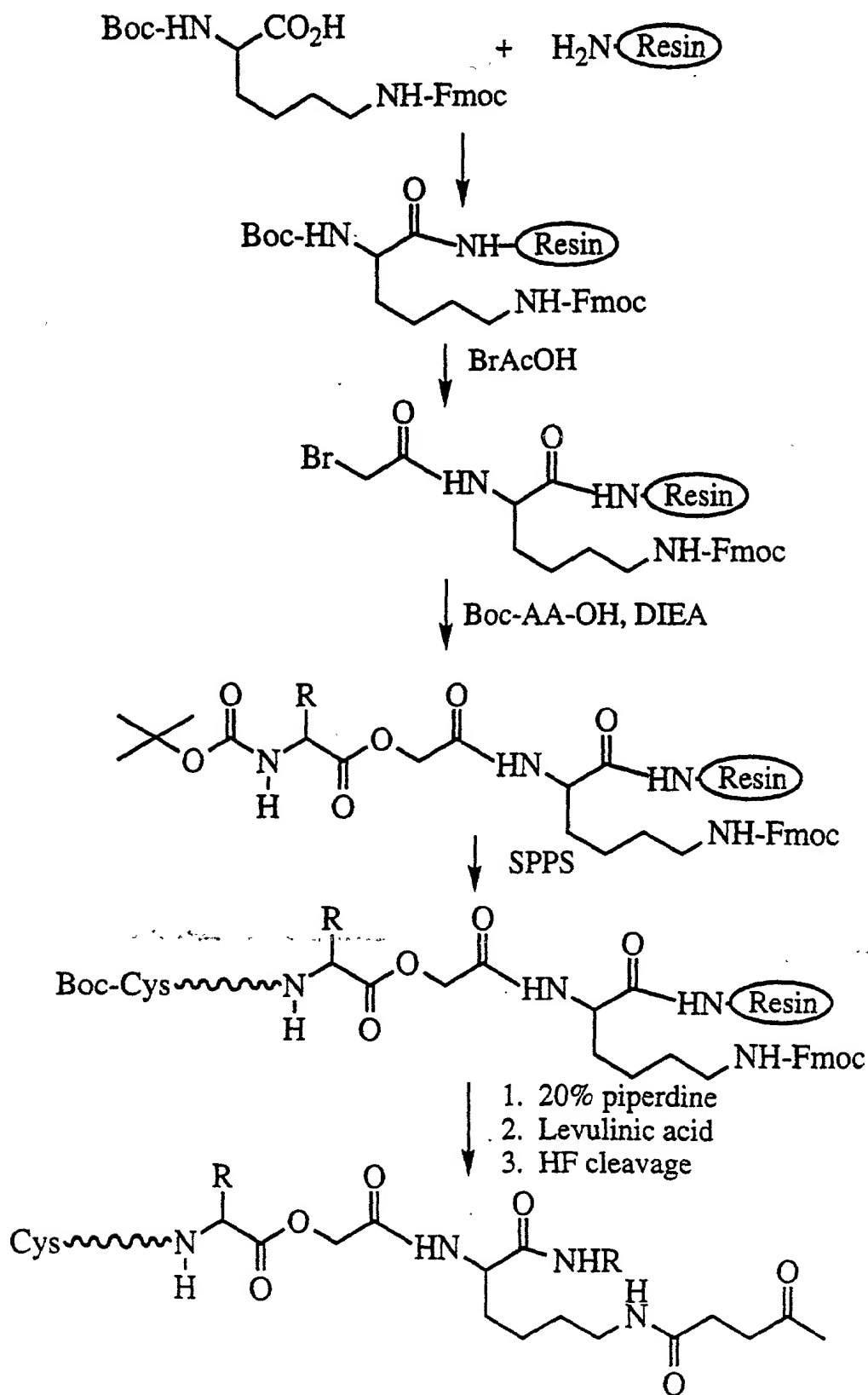


FIG. 27

Synthesis of a Random Sequence by Solid Phase Chemical Ligations in the C- to N-terminal Direction Using Fmoc Protection

ALTKYGFYGCYGRLEEKGCADRKNILA
1 10 19 27

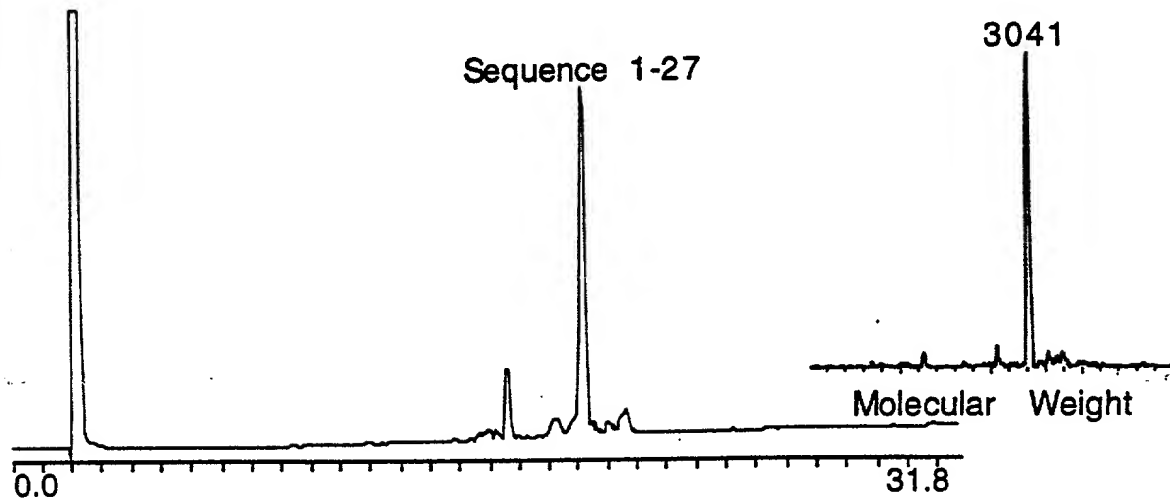


FIG. 28

**Synthesis of a Random Sequence by Solid Phase
Chemical Ligations in the C- to N-terminal Direction
Using ACM Protection**

ALTKYGFYGCYGRLEEKGCADRKNILA
1 10 19 27

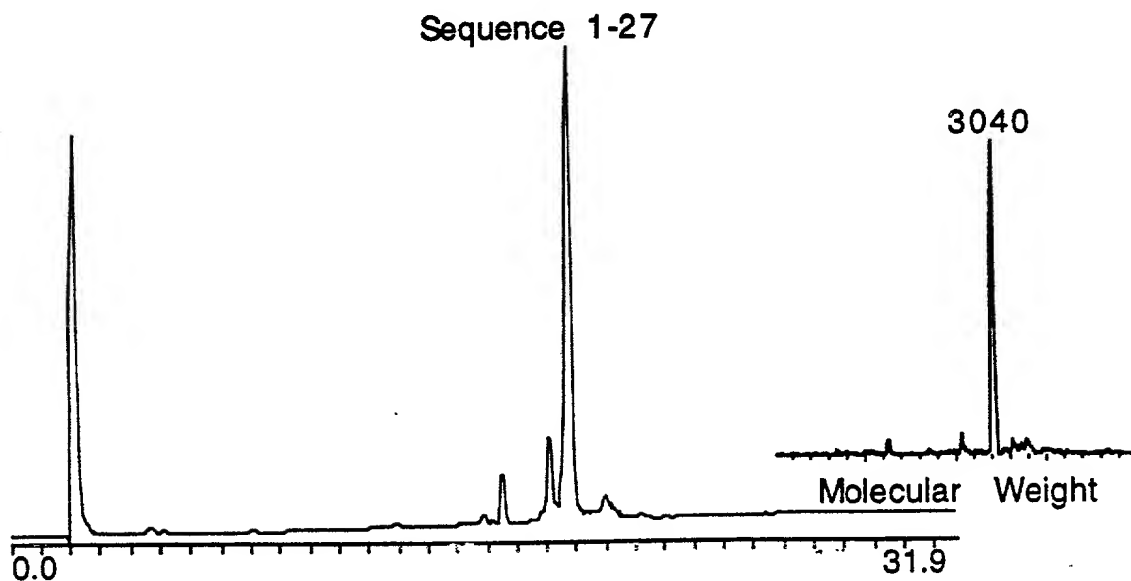


FIG. 29

Synthesis of Phospholipase A2 Group 5 by Solid Phase Chemical Ligations in the C- to N-Terminal Direction

1 26 59
GLLDLKSMEKVTGKNALINYGYG**CYCGNGGRGTPKDGTDWCCWAHDE**CYGRLEEKGC
NIRTQSYKYRFAWGVVTCEPGPFCHVNL**CACDRKL**VYCLRNLR**SYNPQYQYFP**NILCS
88 118

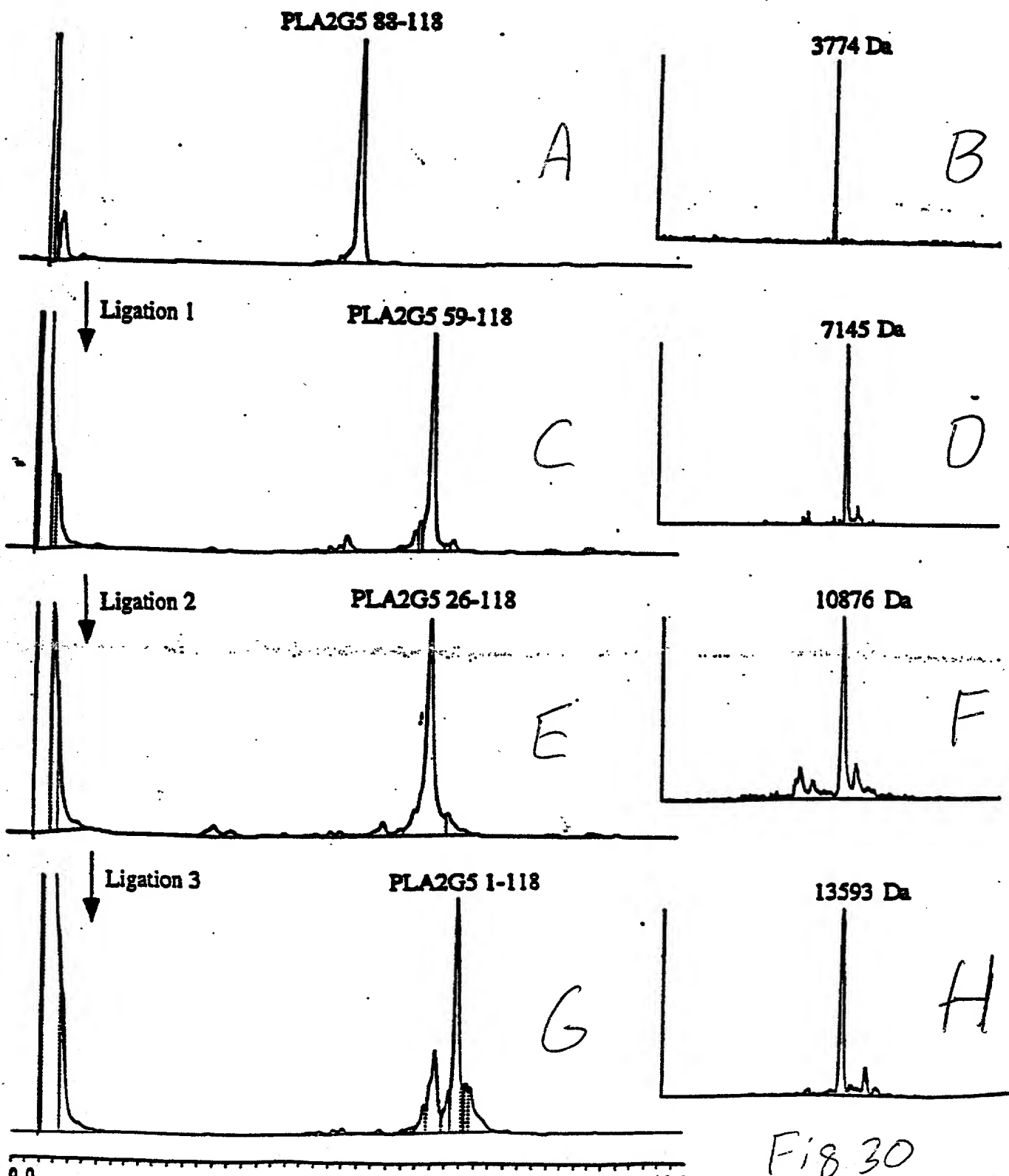
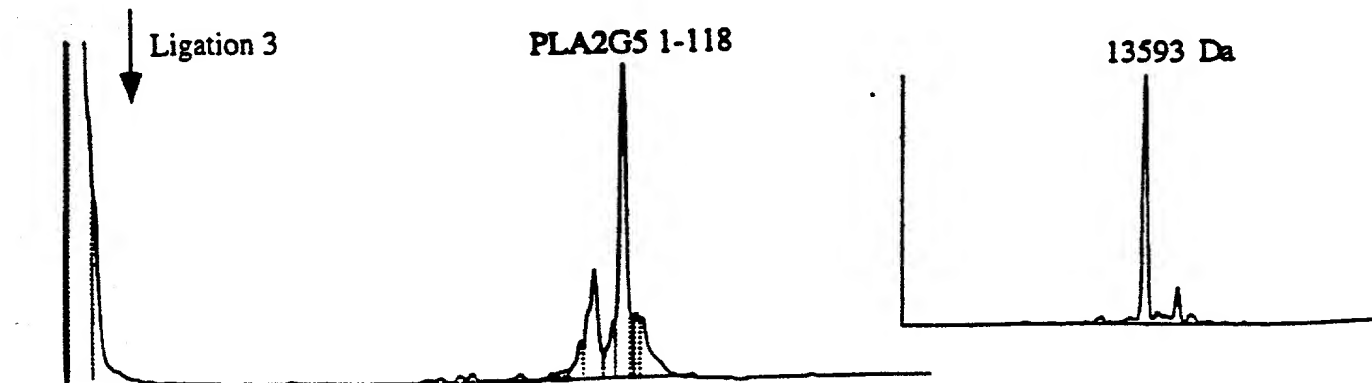
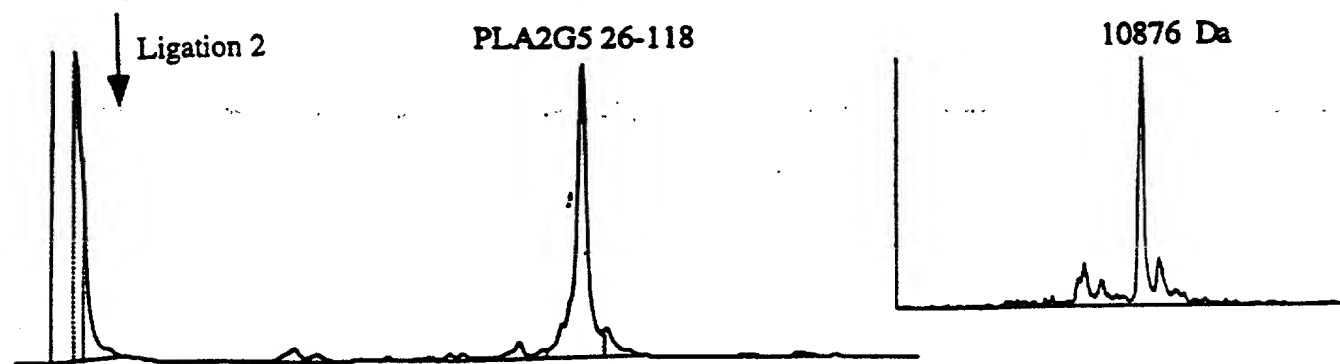
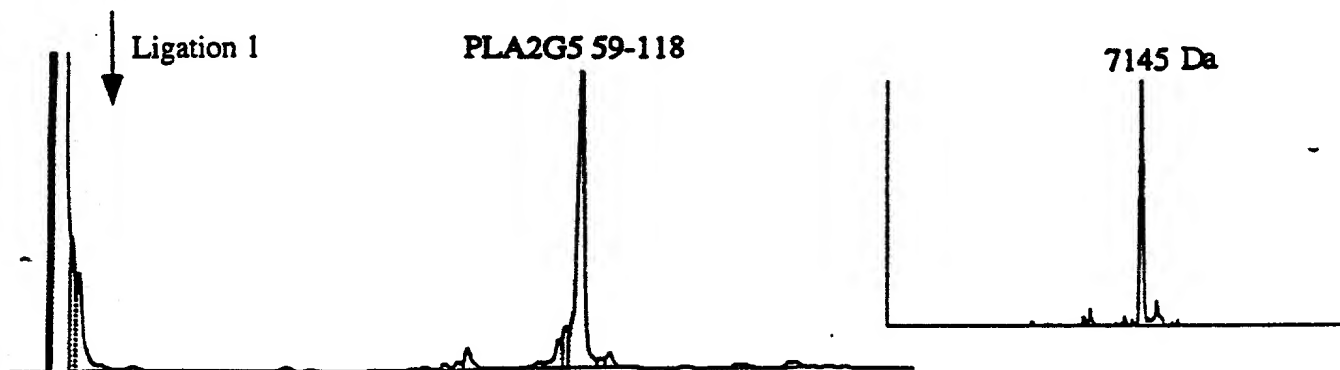
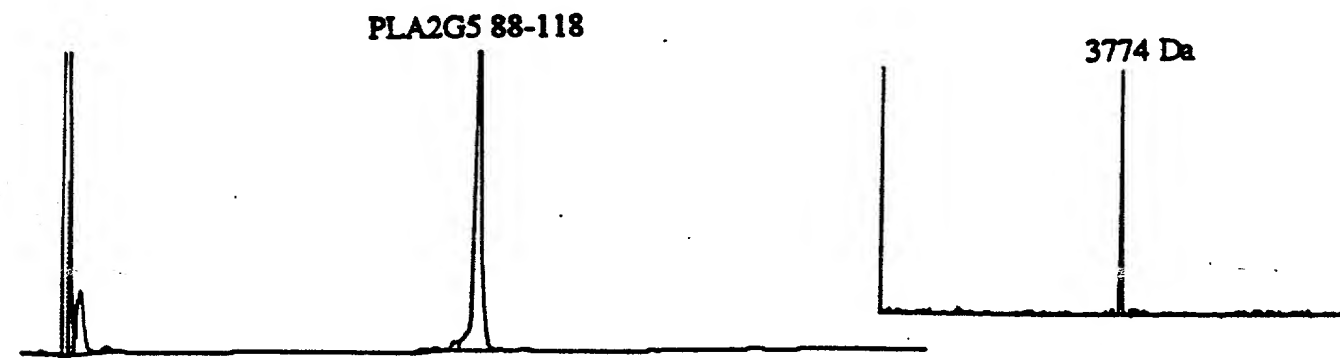


Fig 30

Synthesis of Phospholipase A2 Group 5 by Solid Phase Chemical Ligations in the C- to N-Terminal Direction

1 26 59
 GLLDLKSMIEKVTGKNALTNYGFIYGCYCGWGGRGTPKDGTDWCCWAHDEHCYGRLEERGC
 NIRTQSYKYRFAWGVVTCEPGPFCHVNL**CACDRKLVYCLRNLR**SYNPQYQYFPNILCS
 88 118



0.0

FIG 20

48.9